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(54) Title: COMPOSITION AND METHOD FOR <i>IN VIVO</i> IMAGING OF AMYLOID DEPOSITS (57) Abstract An amyloid binding composition for <i>in vivo</i> imaging of amyloid deposits comprising a labeled amyloid protein or variant thereof which binds to amyloid deposits <i>in vivo</i> ; and a pharmaceutically acceptable carrier, is described. Methods of detecting amyloid deposits and for diagnosing Alzheimer's Disease and Down's Syndrome are also described.		

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COMPOSITION AND METHOD FOR IN VIVO IMAGING OF AMYLOID
DEPOSITS

Background of the Invention

The present invention relates to the identification of compositions which are suitable for use in in vivo imaging of amyloid deposits and methods related thereto.

More specifically, the present invention relates to a method of diagnosing Alzheimer's Disease.

Alzheimer's Disease ("AD") is the most common cause of dementia in the United States, and the presence of the disease is difficult to determine without invasive biopsies. The condition is characterized by impairments in memory, cognition, language and mobility, and these impairments progress over time.

Post-mortem slices of brain tissue from AD victims show that amyloid-containing senile plaques are a prominent feature of selective areas of the AD and the Down Syndrome brain. Divry, P., *J. Neurol. Psych.*, 27: 643-657 (1927); Wisniewski, et al., "Reexamination of the pathogenesis of the senile plaque," In Zimmerman, H.M. (ed.): *Progress in Neuropathology*, N.Y. (1973), Grune and Stratton, pp. 1-26. These plaques range in size from approximately 9 μm to 50 μm in diameter, when viewed by immunocytochemical methods designed to detect amyloid, and they vary in morphology and density. Majocha et al., *Proc. Natl. Acad. Sci. USA*, 85: 6182-6186 (1988). Classical staining methods can detect senile plaques as large as 200 μm . Tomlinson, et al., "Ageing and the dementias," In: Adams, J.H., et al., (ed.), *Greenfield's Neuropathology*, Edition 4, J. Wiley and Sons, N.Y., pp. 951-1006. These plaques are most often found in the cerebral cortex, but they also occur in deeper grey matter, including the amygdaloid nucleus, the corpus striatum, and the diencephalon. Plaques have also been described in the cerebellum. Pro, et al., *Neurology*, 30: 820-825 (1980). Senile plaques are composed of

extracellular amyloid, reactive cells, and degenerating neurites that contain Paired Helical Filaments, lysosomes, abnormal mitochondria and astrocytic processes. Wisniewski, et al., *supra* (1973). The mechanisms responsible for the excessive accumulation of amyloid, the major proteinaceous component of senile plaques, have been recently addressed at the protein chemistry, molecular biology and genetic level.

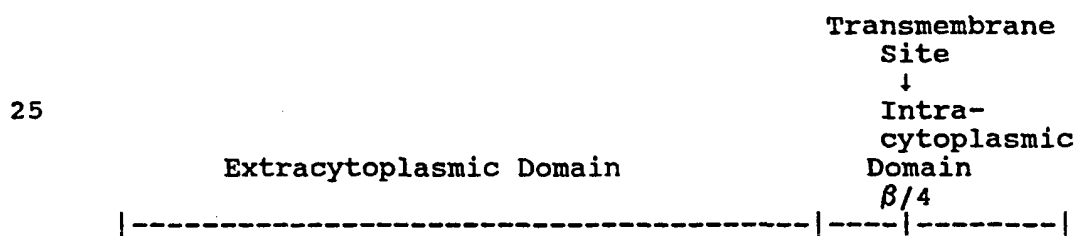
Specifically, amyloid is composed of fibrils of 4-8 nm in diameter that form the core of senile plaques. Mertz et al., *Acta Neuropathol.*, 60: 113-124 (1983). The amyloid is readily demonstrated by application of thioflavin S or Congo red to brain sections. In the latter case, polarized light causes amyloid to appear with a characteristic yellow-green color. The staining property reflects the presence of twisted beta-pleated sheet fibrils, as noted above. A detailed discussion of the biochemistry and histochemistry of amyloid can be found in Glenner, *N. Engl. J. Med.*, 302: 1333-1343 (1980).

Vascular amyloidosis, referred to as congophilic angiopathy, has been recognized since the early part of this century as a significant aspect of the microscopic pathology of Alzheimer's Disease. Vinters, et al. *Stroke*, 18: 311-324 (1987). Over 90% of Alzheimer cases have congophilic angiopathy. Glenner, et al., *Ann. Pathol.*, 1: 120-129 (1981). Similar to parenchymal amyloid deposits, vascular amyloid is demonstrated by characteristic thioflavin S and Congo red staining reactions. The parieto-occipital cortex is usually more affected than that in the frontal and temporal lobes. Tomlinson et al., *supra* (1984).

In vascular amyloidosis, the amyloid appears to infiltrate the micro-vasculature, and affected vessels often pass from the leptomeninges into the cortex. Small cerebral vessels with arterioles that appear as thickened

tubes are observed. The changes include the small pial and intracortical arterioles, the leptomeningeal vessels and the intracortical capillaries. Tomlinson et al., *supra* (1984). Immunocytochemical and electron
 5 microscopic studies have indicated that the amyloid component of senile plaques are often observed in close proximity to affected microvessels. Allsop, et al. *Neurosci. Lett.*, 68: 252-256 (1986). However, the angiopathy may occur without senile plaques. Montjoy, et
 10 al., *J. Neurol. Sci.*, 57: 89-103 (1982).

The principle component of both cerebral (senile plaques) and vascular amyloid is the 4.2 kilodalton peptide, β -amyloid, which is also referred to as β /A4 and A4. Glenner et al., *Biochem. Biophys. Res. Commun.*, 120:
 15 885 (1984). β /A4 is derived from a parent molecule, the amyloid precursor protein (APP). Kang et al., *Nature*, 325: 733-736 (1987). At least three major variants of APP are known, having 695, 751 and 770 amino acids, respectively. In all three variants, the site of the
 20 β /A4 peptide is in the same relative 3'-end location, as follows:



30 Kang et al., *supra* (1987) showed through cloning APP-695 that APP has a large extracellular domain, a transmembrane domain (which gives rise to the β /A4 peptide) and an intracytoplasmic domain (See Figure 9). The signal sequence, for transport through the
 35 endoplasmic reticulum membrane, is followed by a region rich in cysteine, which suggests that disulfide bridges may stabilize this portion of the structure. Within the next 100 residues are a stretch of 7 uninterrupted

threonine residues and a region containing 28 glutamic acid residues and 17 aspartic residues. Marotta, et al. *J. Mol. Neurosci.*, 3: 111-125 (1992) suggest that this domain could bind cations extensively and may have physiological significance. Sodium dodecyl sulfate ("SDS") may be bound to a lesser extent than usual due to this domain. The region from residue 290 to 597, at which point the β /A4 site begins, contains two potential N-glycosylation sites at positions 467-469 and 496-498. The β /A4 peptide (residues 596-638 or 639) is either 42 or 43 amino acids in length and partly includes the putative transmembrane domain (amino acids 625-648). The C-terminal region of the APP is relatively small, consisting of 57 residues.

Following the transmembrane region, lysine residues are present (residues 649-651) which, according to Kang et al. *supra* (1987), could interact with phospholipid head groups in the membrane. This feature has been described for the junction between membrane and cytoplasmic domains of cell-surface receptors. One site (amino acids 684-686) is a potential glycosylation sequence.

Gandy et al. report that during in vitro studies of synthetic peptides corresponding to the cytoplasmic domain, it was observed that protein kinase C rapidly catalyzed the phosphorylation of a peptide corresponding to amino acid residues 645-661 on ser-655. Gandy et al., *Proc. Natl. Acad. Sci. USA*, 85(16): 5218-5221 (1988), suggesting that this site may be an important control region for amyloid metabolism and its interaction with other intracellular regulatory elements.

Recent research has also focused on the biological activity of β /A4. Specifically, it has been noted that this peptide and its fragments are trophic, toxic and or amnestic at various concentrations. Also, β /A4 forms insoluble aggregates (self-aggregates) under various

conditions, and the neurotoxicity of β /A4 is related to the aggregation process. Kirshner, et al., *Proc. Natl. Acad. Sci. USA*, 84: 6953-6957 (1987) and Maggio, *Annu. Rev. Neurosci.*, 11: 13-28 (1988). Maggio et al., also
5 studied the aggregation properties of radioiodinated synthetic β /A4 peptides *in vitro*. *Proc. Natl. Acad. Sci. USA*, 89: 5462-5466 (June 1992).

Pike et al., *J. Neurosci.*, 13(4): 1676-1687 (1993) tested the aggregation properties of an overlapping
10 series of synthetic β -amyloid peptides and compared them with their neurotoxic properties *in vitro*. They discovered that few peptides assembled into aggregates immediately after solubilization but that over time peptides containing the highly hydrophobic 829-35 region
15 formed stable aggregations. In short-term cultures, neurotoxicity was associated with those peptides demonstrating significant aggregations.

Thus far, diagnosis of AD has been achieved mostly through clinical criteria evaluation, brain biopsies and
20 post mortem tissue studies. However, recent work has focused on immunoassay methods for detecting markers of AD in body fluids such as spinal fluid and also in *in situ* hybridization studies using nucleic acid probes. World Patent No. 92/17152 by Potter; Warner, M., *Anal. Chem.*, 59: 1203A (1987); U.S. Patent No. 4,666,829 by
25 Glenner et al. In U.S. application no. 105,751, the contents of which is hereby incorporated by reference, Marotta et al. describe anti- β /A4 antibodies for purposes of *in vitro* and *in vivo* diagnostic methods.

30 Glenner et al., *supra*, teach the use of the β /A4 peptide, or fragments thereof, for the production of antibodies which recognize the antigenic determinants of the polypeptide or homologues thereof. Glenner et al. further teach the use of the disclosed polypeptide for
35 the production of nucleic acid probes which hybridize with the gene encoding the polypeptide. One such

polypeptide has the following amino acid sequence (SEQ ID NO:1): H₂N-Asp-Ala-Glu-Phe-Arg-His-Asp-Ser-Gly-Tyr-Gln-Val-His-His-Gln-Lys-Leu-Val-Phe-Phe-Ala-Glu-Asp-Val-Gly-Ser-Asn-Lys-COOH. The diagnostic methods taught in this

5 patent are characterized as non-invasive.

U.S. Patent Nos. 5,039,511 and 4,933,156 by Quay et al. describe the *in vitro* and *in vivo* use of iodinated imaging compounds derived from bisdiazobenzidine compounds to detect the presence and location of amyloid

10 deposits in an organ or area of a patient.

Although β /A4 has been considered for use in *in vitro* diagnostic methods, this polypeptide has never been described in connection with *in vivo* diagnostic imaging methods. Therefore, a need exists for a diagnostic

15 *in vivo* imaging method that exploits the self-aggregation properties of amyloid proteins such as β /A4.

Summary of the Invention

One object of the present invention is to provide an amyloid binding composition for *in vivo* imaging of amyloid deposits comprising a labeled amyloid protein which binds to amyloid deposits *in vivo* and a pharmaceutically acceptable carrier.

Another object of the present invention is to provide an *in vivo* method for detecting amyloid deposits in a subject comprising the steps of administering to a subject a detectable quantity of an amyloid binding composition comprising a labeled amyloid binding protein and a pharmaceutically acceptable carrier and detecting the binding of the labeled protein to the amyloid deposit.

Another object of the present invention is to provide a method of diagnosing an amyloidosis-associated disease, such as Alzheimer's Disease and Down Syndrome, by applying the above method to the detection of amyloid deposits in subjects suspected of having an amyloidosis-associated disease.

The amyloid binding protein of the present invention includes all variants of the amyloid protein which bind to amyloid deposits *in vivo*.

Brief Description of the Drawings

Figure 1 shows a chart setting forth chemically documented amyloidosis with protein types.

Figure 2 depicts a PAGE-SDS gel of the A4-O synthetic amyloid peptide. The amyloid polypeptide of 28 residues, corresponding to the previously reported sequence of Masters et al., *Proc. Natl. Acad. Sci. USA* 82: 4245-4249 (1985) was synthesized on a Biosearch SAM2 synthesizer using the general procedure of Merrifield, *J. Am. Chem. Soc.*, 85: 2149-2154 (1963). Purification was achieved with a 3 X 65 cm column of Sephadex G50 (10-40 μ). The peptide (10 μ g) was suspended in sample buffer containing 2% SDS (Brown, et al., *J. Neurochem.*, 40: 299-308 (1983))

and 9.5 M urea. Electrophoresis was carried out on a uniform 10% gel containing 0.1% SDS. (A) Molecular weight markers: phosphorylase B (94 kd), bovine serum albumin (68 kd), ovalbumin (43 kd), carbonic anhydrase (29 kd), trypsin inhibitor (22 kd), lysozyme (14.5 kd). (B) The synthetic amyloid peptide ran as a sharp band at the front of the gel and as an aggregated form of higher molecular weight.

Figure 3 depicts gel electrophoresis of the synthetic peptides A4-O and P2 (APP amino acids 413-429). (Panel a): Twenty ug of A4-O (lane 1) and P2 (APP amino acids 413-429) (lane 2) were analyzed by 18% SDS-PAGE (acrylamide : bisacrylamide = 30 : 0.8). (Panel b): Analysis of A4-O and P2 (APP amino acids 413-429) on 11% SDS/urea-PAGE (acrylamide : bisacrylamide = 20 : 1). Lanes 1-3 containing A4-O (10 ug) were incubated with 2% SDS and 5% 2-ME at 95°C for 5 minutes (Lane 1), 30 min (Lane 2) and 60 minutes (Lane 3). Lane 4 contained P2 (10ug). Gels were stained with Coomassie Brilliant blue R-250. Molecular weights are shown on the right (Kd).

Figure 4 shows slot blots of immunostained A4-O after addition of itself or a second A4 homologue. This assay depicts the increase staining intensity after A4 homologues are added to one another. This reflects the ability of homologues to self-aggregate and thus increase the staining intensity. In each case, the slot contained 1 ug of A4 peptide. To each, 2.5, or 10.0 ug of exogenous peptide was added, as indicated. The blots were then immunostained (see descriptions of Figures 5, 6 and 7).

Figure 5 shows immunoblots with and without exogenous A4-O peptide. Density values of the immunoreaction products of A4-O with and without exogenous peptides after reaction with 10H3. The values of the bars correspond to the density of blots shown in Figure 4. The description of Figure 4 indicates the condition of

each blot with regard to the exogenous peptide that was added to the blotted peptide prior to addition of 10H3. The height of the bars above the black bar (no peptide addition) is a measure of the extent to which the exogenous peptide bound to the attached peptide on the filter paper and increased the density of immunostaining of the complex.

At the three indicated concentrations (2.5, 5.0 and 10.0 ug/ml) A4-0 was added to the A4-0 that was already present at a concentration of 1 ug. The blot was then immunostained to develop the colored reaction product. The data were derived from scanning Figure 4.

Figure 6 shows immunoblots with and without exogenous A4-H peptide. At the three indicated concentrations (2.5, 5.0 and 10.0 ug/ml) A4-H as added to the A4-0 that was already present at a concentration of 1 ug. The blot was then immunostained to develop the colored reaction product. The data were derived from scanning Figure 4. See description of Figure 5 for more details.

Figure 7 shows immunoblots with and without exogenous Op1 peptide. At the three indicated concentrations (2.5, 5.0 and 10.0 ug/ml) Op1 as added to the A4-0 that was already present at a concentration of 1 ug. The blot was then immunostained to develop the colored reaction product. The data were derived from scanning Figure 4. See description of Figure 5 for more details.

Figure 8 shows the reactivity of 10H3 towards A4-0 (upper panel).

Figure 9 (SEQ ID NOS:11 and 12) is the nucleotide sequence and predicted amino acid sequence of cDNA encoding the precursor protein (APP) of the β /A4 with the β /A4 region boxed, as set forth in Kang et al., *supra*, (1987).

Detailed Description of the Preferred Embodiments

Applicants have discovered that an amyloid binding composition comprising a labeled amyloid protein may be used *in vivo* for detecting the presence and location of amyloid deposits. The amyloid binding composition of the present invention comprises a labeled amyloid protein and a pharmaceutically acceptable carrier. This protein is any natural or synthetic protein which binds to amyloid deposits *in vivo*. In one embodiment, the protein is the β -amyloid polypeptide (B/A4 peptide), which in its longest form has 42 to 43 amino acid residues, as shown in Figure 9. See Masters, et al., *Proc. Nat. Acad. Sci., USA.*, 82: 4245-4249 (1985).

As noted above, B/A4 is derived from a larger amyloid precursor protein having from 695 to 770 amino acids. See Kang et al., *Nature*, 325: 733 (1987). The term "amyloid deposit" includes amorphous, eosinophilic materials that are associated with amyloidosis, a disease complex including over 20 different clinically defined syndromes, as discussed above. Chemically, amyloid deposits are proteinaceous, and their chemical compositions are unique for each of the clinical syndromes with which they are associated, as set forth in Figure 1. Preferably, the amyloid deposit of the present invention is that found in the brain of Alzheimer's Disease patients. As noted above, such amyloid deposits are found in senile plaques in selected areas of the AD brain and are composed of fibrils of 4-8 nm diameter. These plaques are detected by application of thioflavin S or Congo red to brain sections and in the latter case, appear yellow-green under polarized light. They have twisted beta-pleated sheet fibrils and are further characterized by Glenner, *N. Eng. J. Med.*, 302: 1333-1343 (1980). In another embodiment, the amyloid deposit of the present invention is that which is associated with vascular amyloidosis, as

-11-

described in Vinters, *Stroke*, 18: 311-324 (1987). Vascular amyloid deposits infiltrate the cerebral microvasculature. Similar to amyloid deposits found in senile plaques in the parenchyma of the AD brain, vascular amyloid deposits have characteristic thioflavin S and Congo red staining reactions. Montjoy et al., *J. Neurol. Sci.*, 57: 89-103 (1988).

The term "amyloidosis-associated disease" includes any disease characterized by local or systemic amyloid deposits. (See Figure 1) Preferably, the amyloidosis-associated disease of the present invention is Alzheimer's Disease or Down Syndrome.

In addition to amyloid protein purified from natural sources such as cerebrovascular tissue, as described hereinafter, amyloid protein of the present invention includes recombinant and synthetic amyloid protein and variants of the naturally occurring, recombinant and synthetic protein. In a preferred embodiment, the amyloid protein of the invention comprises the β -amyloid polypeptide and variants thereof. The category of "variants" includes, for example, a fragment of the β -amyloid polypeptide or any homologous amino acid sequence or amino acid addition, wherein the resulting polypeptide has the same or similar function as the natural occurring polypeptide in that it binds to amyloid deposits *in vivo*. In one embodiment, the amyloid protein of the present invention is comprised of the β -amyloid polypeptide or variant thereof and amino acids from the APP protein which are from regions of the APP protein which are either adjacent or non-adjacent to the β -amyloid polypeptide. For example, in one embodiment, the amyloid protein of the present invention comprises:

(A) The β /A4 peptide alone (SEQ ID NO:2):

Asp-Ala-Glu-Phe-Arg-His-Asp-Ser-Gly-Tyr-Glu-Val-His-His-
Gln-Lys-Leu-Val-Phe-Phe-Ala-Glu-Asp-Val-Gly-Ser-Asn-Lys-

-12-

Gly-Ala-Ile-Ile-Gly-Leu-Met-Val-Gly-Gly-Val-Val-Ile-Ala-Thr; or

(B) The β /A4 peptide plus the amino acids of the transmembrane domain of the APP (SEQ ID NO:3):
5 Asp-Ala-Glu-Phe-Arg-His-Asp-Ser-Gly-Tyr-Glu-Val-His-His-Gln-Lys-Leu-Val-Phe-Phe-Ala-Glu-Asp-Val-Gly-Ser-Asn-Lys-Gly-Ala-Ile-Ile-Gly-Leu-Met-Val-Gly-Gly-Val-Val-Ile-Ala-Thr-Val-Ile-Val-Ile-Thr-Leu-Val-Met-Leu; or

(C) The β /A4 peptide plus the remaining C-terminal amino acids of the entire APP (SEQ ID NO:4):
10 Asp-Ala-Glu-Phe-Arg-His-Asp-Ser-Gly-Tyr-Glu-Val-His-His-Gln-Lys-Leu-Val-Phe-Phe-Ala-Glu-Asp-Val-Gly-Ser-Asn-Lys-Gly-Ala-Ile-Ile-Gly-Leu-Met-Val-Gly-Gly-Val-Val-Ile-Ala-Thr-Val-Ile-Val-Ile-Thr-Leu-Val-Met-Leu-Lys-Lys-Lys-Gln-15 Tyr-Thr-Ser-Ile-His-His-Gly-Val-Val-Glu-Val-Asp-Ala-Ala-Val-Thr-Pro-Glu-Glu-Arg-His-Leu-Ser-Lys-Met-Gln-Gln-Asn-Gly-Tyr-Glu-Asn-Pro-Thr-Tyr-Lys-Phe-Phe-Glu-Gln-Met-Gln-Asn; or

(D) The β /A4 peptide with the preceeding 10 amino acids of the APP (SEQ ID NO:5):
20 Lys-Thr-Glu-Glu-Ile-Ser-Glu-Val-Lys-Met-Asp-Ala-Glu-Phe-Arg-His-Asp-Ser-Gly-Tyr-Glu-Val-His-His-Gln-Lys-Leu-Val-Phe-Phe-Ala-Glu-Asp-Val-Gly-Ser-Asn-Lys-Gly-Ala-Ile-Ile-Gly-Leu-Met-Val-Gly-Gly-Val-Val-Ile-Ala-Thr; or

(E) The β /A4 peptide with any other APP amino acids attached to it that are not normally adjacent (SEQ ID NO:6):
25

X-Asp-Ala-Glu-Phe-Arg-His-Asp-Ser-Gly-Tyr-Glu-Val-His-His-Gln-Lys-Leu-Val-Phe-Phe-Ala-Glu-Asp-Val-Gly-Ser-Asn-30 Lys-Gly-Ala-Ile-Ile-Gly-Leu-Met-Val-Gly-Gly-Val-Val-Ile-Ala-Thr-Y,

wherein X and Y are one or more APP amino acids which are not adjacent to β /A4 in the nature; and

(F) any fragment of (A)-(E), wherein said fragment is large enough to bind amyloid deposit
35 in vivo.

The term "fragment" includes a linear amino acid subsequence of the β -amyloid polypeptide, wherein such fragment binds amyloid deposits *in vivo*. A variant which contains an amino acid sequence variation or substitution is a homologous sequence. "Homology" between two sequences connotes a likeness short of identity indicative of a derivation of the first sequence from the second. For example, a polypeptide is "homologous" to β -amyloid polypeptide if it contains an amino acid sequence similar enough to the natural sequence so as to confer the same or similar amyloid binding property as the natural β -amyloid polypeptide. Such a sequence may be only a few amino acids long and may be a single linear sequence or one or more linear sequences which confer binding activity to the polypeptide when amino acids from separated portions of a linear sequence are spatially juxtaposed after protein folding. The variants encompassed by this invention can be ascertained, for example, by the *in vitro* quantitative assays describe below in Examples 3-7. That is, applicants have conducted a series of studies involving the addition of increasing concentrations of β -amyloid polypeptide variants to a solid support containing a specific peptide called A4-0. The increase in density of immunostain using an anti-A4-0 monoclonal antibody, 10H3, described in U.S. Patent application no. 105,751 by Marotta, et al., incorporated by reference above, was measured. Based upon this work, it was possible to determine which peptides were suitable for use in the *in vivo* methods, according to the invention. Other poly- and/or monoclonal antibodies suitable for this assay can be produced by methods well known in the art. See Kennett et al., *Monoclonal Antibodies- Hybridomas: A New Dimension in Biological Analysis*, Plenum Press (1980)

Protein which qualifies as "amyloid protein" according to the above criteria can be produced by

methods known and emerging in the art, including conventional reverse genetic techniques, i.e., by designing a genetic sequence based upon an amino acid sequence or by conventional genetic splicing techniques.

5 For example, β -amyloid polypeptide variants can be produced by techniques which involve site-directed mutagenesis or oligonucleotide-directed mutagenesis. See, for example, "Mutagenesis of Cloned DNA," in CURRENT PROTOCOLS IN MOLECULAR BIOLOGY 8.0.3 et seq. (Ausubel, et

10 al. eds. 1989) ("Ausubel").

Other amyloid protein variants within the present invention are molecules that correspond to a portion of the β -amyloid polypeptide, but are not coincident with the natural molecule, and display the binding activity of

15 the natural molecule when presented alone or, alternatively, when linked to a carrier or biologically active signal sequence that permits proteins to pass through membranes. See von Heijne, G., J. Mol. Biol., 184: 99-105 (1985). An amyloid protein variant of this

20 type could represent an actual fragment, as discussed above, or could be a polypeptide synthesized *de novo* or recombinantly.

To be used in recombinant expression of amyloid protein or amyloid protein variant, a polynucleotide

25 molecule encoding such a molecule would preferably comprise a nucleotide sequence, corresponding to the desired amino acid sequence, that is optimized for the host of choice in terms of codon usage, initiation of translation, and expression of commercially useful

30 amounts of, for instance, β -amyloid polypeptide or β -amyloid polypeptide variant. Also, the vector selected for transforming the chosen host organism with such a polynucleotide molecule should allow for efficient maintenance and transcription of the sequence encoding

35 the polypeptide. The encoding polynucleotide molecule may code for a chimeric protein; that is, it can have a

nucleotide sequence encoding a biologically active part of the β -amyloid molecule operably linked to a coding sequence for a non- β -amyloid moiety, such as a signal peptide for the host cell.

5 For instance, in order to isolate a DNA segment which encodes a β -amyloid molecule, total DNA from cerebrovascular tissue can be prepared according to published methods. See, for example, Maniatis, et al., MOLECULAR CLONING: A LABORATORY MANUAL (Cold Spring
10 Harbor Laboratories, NY 1982); Baess, *Acta Pathol. Microbiol. Scand. (Sect. B)*, 82: 78084 (1974). The DNA thus obtained can be partially digested with a restriction enzyme to provide an assortment of genomic fragments. An enzyme with a tetranucleotide recognition
15 site, such as *Sau3A (MboI)*, is suitable for this purpose. The fragments from such a partial digestion then can be size-fractionated, for example, by sucrose gradient centrifugation (see Maniatis, *supra*) or by pulsed field gel electrophoresis (See Anad, *Trends in Genetics*,
20 November 1986, at pages 278-83), to provide fragments of a length commensurate with that of DNA encoding the β -amyloid molecule. Molecular cloning of amyloid cDNA derived from mRNA of the Alzheimer brain and the expression thereof is described in detail in Zain et al.,
25 *Proc. Natl. Acad. Sci. USA.*, 85: 929-933 (1988) and Marotta et al., *Proc. Natl. Acad. Sci. USA.*, 86: 337-341 (1989), respectively, both of which are herein incorporated by reference.

 According to well-known methods described, for
30 example, in Ausubel at 5.0.1 et seq., the selected fragments can be cloned into a suitable cloning vector. A DNA sequence thus obtained could be inserted, for example, at the *BamHI* site of the pUC18 cloning vector which is transfected into appropriate host cells such as
35 *E. coli* or a mammalian cell. A variety of screening

mechanisms known in the art of the invention can then be used to identify clones containing the β -amyloid gene.

In another embodiment of the invention, amyloid protein of the present invention is purified from tissue samples. For example, brains from Alzheimer's Disease post-mortum patients are histologically sectioned and stained with Congo Red dye. Upon visualization with a polarizing microscope, amyloid deposits can be identified by their green color. Brains exhibiting extensive cerebrovascular amyloidosis are used as source for purified amyloid protein. After removal of contaminants from the amyloid containing vessels of the meninges, the meningeal tissues are homogenized and centrifuged to yield a brownish layer rich in amyloid fibrils. This layer is then digested with collagenase, solubilized in 6M guanidine HCl, pH 8.0 and centrifuged. The supernatant containing the solubilized protein is desalted by dialysis and gel exclusion column chromatography and high performance liquid chromatography is used to purify the polypeptide. The amino acids for the purified protein (e.g., β -amyloid polypeptide) are then sequentially cleaved in an automated amino acid sequencer, such as a Beckman 890 C spinning cup sequencer, and analyzed by high performance liquid chromatography in order to determine the amino acid sequence of the amyloid protein. See Glenner & Wong, *Biochem. Biophys. Chem. Res. Commun.*, 120: 885 (1984).

In another embodiment, amyloid protein and variants thereof can be produced in accordance with published methods. For instance, Kirschner et al., *Proc. Natl. Acad. Sci. USA*, 84: 6953-57 (1987) used an ABI Synthesizer model 380 B (Applied Biosystems, Foster City, CA) to synthesize synthetic β -amyloid peptides consisting of residues 1-28 and homologues thereof. General methods for peptide synthesis can be found in Clark Lewis et al., *Science*, 231: 134 (1986). See also, Hilbich et al., *J.*

Mol. Biol., 218: 149-163 (1991); Majocha et al., *Proc. Natl. Acad. Sci. USA*, 85: 6182-6186 (1988); and U.S. Patent application No. 105,751 by Marotta et al.

5 The term "in vivo imaging" refers to any method that permits the detection of a labeled amyloid protein which binds to amyloid deposits located in a subject's body. A "subject" is a mammal, preferably a human. Often, particularly when the composition and method of the invention is directed to the diagnosis of Alzheimer's
10 Disease or Down Syndrome, the subject will manifest clinical symptoms of the suspected amyloidosis. These clinical symptoms are well-known to the practitioner of this invention and include loss of memory, and other impairments described above.

15 The amyloid binding composition of the present invention must be of a "detectable quantity." A detectable quantity is that which is sufficient to enable detection of the site of amyloid deposit location when compared to a background signal. The dosage of the
20 amyloid binding composition will vary depending upon such considerations as age, condition, sex, extent of disease in the patient, counterindications, and other variables, to be adjusted by the individual physician. Dosage can vary from
25 0.01 mg/kg to 2,0000 mg/kg, preferably 0.1 mg/kg to 1,000 mg/kg.

In accordance with this invention, the amyloid protein may be labeled by any of several techniques known to the art. See, e.g., Wagner et al., *J. Nucl. Med.*, 20:
30 428 (1979); Sundberg et al., *J. Med. Chem.*, 17: 1340 (1974) and Saha et al., *J. Nucl. Med.*, 6: 542 (1976).

The label is chosen based upon the type of detection instrument employed. For instance, a chosen radionucleotide must have a type of decay which is
35 detectable for a given type of instrument. Another

consideration relates to the half-life of the isotope. The half-life should be long enough so that it is still detectable at the time of maximum uptake by the target, but short enough so that the host does not sustain deleterious radiation. Preferably, the chosen label will lack a particulate emission, but will produce a large number of photons in a 140-200 keV range, which may be readily detected by, for instance, conventional gamma camera. Suitable radioisotopes for purposes of this invention include, gamma-emitters, position-emitters, x-ray emitters and fluorescence-emitters. These radioisotopes include Iodine-131, Iodine-123, Iodine-126, Iodine-133, Bromine- 77, Indium-111, Indium-113m, Gallium-67, Gallium-68, Ruthenium-95, Ruthenium-97, Ruthenium-103, Ruthenium-105, Mercury-107, Mercury-203, Rhenium-99m, Rhenium-105, Rhenium 101, Tellurium-121m, Tellurium-122m, Tellurium-125m, Thulium-165, Thulium-167, Thulium-168, Technetium-99m and Fluorine-18. The preferred radiolabel is Technetium-99m. Suitable paramagnetic isotopes for use in Magnetic Resonance Imaging (MRI), according to this invention, include ^{157}Gd , ^{55}Mn , ^{162}Dy , ^{52}Cr , and ^{56}Fe .

Administration to the subject may be accomplished intraventricularly, intravenously, intraarterially, via the spinal fluid or the like. Administration may also be intradermal or intracavitary, depending upon the body site under examination. After a sufficient time has lapsed for the labeled amyloid protein to bind with amyloid deposits, for example 30 minutes to 48 hours, the area of the subject under diagnosis is examined by routine imaging techniques such as MRI, SPECT and planar scintillation imaging. The exact protocol will necessarily vary depending upon factors specific to the patient, as noted above, and depending upon the body site under examination, method of administration and type of label used; the determination of specific procedures

would be routine to the skilled artisan. The distribution of the bound radioactive isotope and its decrease with time is then monitored and recorded. By comparing the results with data obtained from studies of clinically normal individuals, the presence and location of amyloid deposits can be determined.

Thus, in one embodiment, the methods of the present invention is used to diagnoses an amyloidosis-associated disease. Where the site of examination is the brain, the *in vivo* detection of amyloid deposits according to the methods of the present invention signifies a diagnosis of Alzheimer's Disease. The detection of amyloid deposits in the brain of patients manifesting clinical symptoms of Down Syndrome, signifies a diagnosis of Down Syndrome. In that regard, applicants note that the gene for APP, located on chromosome 21, is over-represented in Down Syndrome individuals (Serra et al., *Amer. J. Med. Gen. Supp.*, 7: 11-19 (1990)). Accumulations of amyloid occur in young Down Syndrome patients, with nearly 90% of Down Syndrome subjects aged less than 30 years showing amyloid accumulation (Hyman, *Prog. Clin. Biol. Res.* 379: 123-142 (1992)). The Down Syndrome patient displays amyloid accumulations early in life, often by late teenage years. As adults, nearly 100% will develop Alzheimer Disease (Cork, *Amer. J. Med. Gen. Supp.*, 7: 282-539 (1990)). The neuropathology of Down Syndrome is essentially identical to that of Alzheimer Disease and includes β /A4 amyloid deposits in senile plaques. The Alzheimer - like lesions represent a major neuropathologic trait of the brain of the Down Syndrome patient (Serra et al., *Supra* (1990)).

The amyloid-binding compositions of the present invention are advantageously administered in the form of injectable compositions. A typical composition for such purpose comprises a pharmaceutically acceptable carrier. For instance, the composition may contain about 10 mg of

human serum albumin and from about 20 to 200 micrograms of the labeled amyloid protein per milliliter of phosphate buffer containing NaCl. Other pharmaceutically acceptable carriers include aqueous solutions, non-toxic excipients, including salts, preservatives, buffers and the like, as described in REMINGTON'S PHARMACEUTICAL SCIENCES, 15th Ed. Easton: Mack Publishing Co. pp 1405-1412 and 1461-1487 (1975) and THE NATIONAL FORMULARY XIV., 14th Ed. Washington: American Pharmaceutical Association (1975), the contents of which are hereby incorporated by reference. Examples of non-aqueous solvents are propylene glycol, polyethylene glycol, vegetable oil and injectable organic esters such as ethyloleate. Aqueous carriers include water, alcoholic/aqueous solutions, saline solutions, parenteral vehicles such as sodium chloride, Ringer's dextrose, etc. Intravenous vehicles include fluid and nutrient replenishers. Preservatives include antimicrobials, anti-oxidants, chelating agents and inert gases. The pH and exact concentration of the various components of the binding composition are adjusted according to routine skills in the art. See GOODMAN AND GILMAN'S THE PHARMACOLOGICAL BASIS FOR THERAPEUTICS (7th ed.). The skilled artisan would also readily appreciate that a suitable excipient or carrier would need to prevent aggregation of the binding composition prior to contacting the target amyloid deposit *in vivo*.

Particularly preferred amyloid binding compositions of the present invention are those that, in addition to binding to amyloid deposits *in vivo*, are also non-toxic at appropriate dosage levels, have a satisfactory duration of effect, and display an adequate ability to cross the blood-brain barrier. In this regard, United States Patent No. 4,540,564 discloses an approach for enhancing blood-brain barrier-penetrating ability by attaching a centrally acting drug species to a reduced,

biooxidizable, lipoidal form of dihydropyridine pyridinium salt redox carrier. Thus, in one embodiment, the composition of the present invention includes such a blood-brain barrier crossing enhancer carrier.

5 *In vivo* animal testing provides yet a further basis for determining dosage ranges, efficacy of transfer through the blood barrier and binding ability. Particularly preferred for this purpose is the "senile animal" model for cerebral amyloidosis -- animals such as
10 aged dogs or monkeys, which are known to develop variable numbers of Alzheimer-type cerebral senile plaques, see Wisniewski, et al., *J. Neuropathol. & Exp. Neurol.*, 32: 566 (1973), Selkoe, et al., *Science*, 235: 873 (1987) are tested for binding and detection efficacy. This *in vivo*
15 assay requires control-biopsy monitoring to confirm and quantify the presence of amyloid deposits.

Also, cellular models of amyloidosis have been prepared that overproduce β -amyloid polypeptide in animals for purposes of testing the efficacy of the
20 amyloid binding compositions and methods of the present invention. See Marotta, et al. *Proc. Natl. Acad. Sci. USA*, 86: 337-341 (1989). Such cell models have been adapted to a behavior paradigm. See Tate-Ostroff, *Proc. Natl. Acad. Sci. USA* 89: 7090-7094, (1992). That is,
25 because AD patients suffer circadian rhythm dysfunction, this behavioral deficit was modeled in rats by a cell grafting techniques. PC12 cells transfected with the β -amyloid polypeptide C-terminal region of the APP were implanted into the suprachiasmatic nuclei ("SCN") of
30 rats; the SCN is a primary circadian oscillator in mammals. Animals receiving amyloidotic cell grafts, but not animals receiving control cell grafts, exhibited disrupted activity rhythms, although temperature rhythms were unaffected. The specificity of the disruption was
35 similar to circadian dysfunction seen in AD patients. The data supported an association between a defined

behavioral disruption and amyloid overexpression either directly or through the release of cellular factors as a consequence of amyloid overproduction.

5 Other suitable animal models for use in testing the compositions and methods of the present invention are produced transgenically. For instance, Quon et al., *Nature*, 352: 239-241 (1991) used rat neural-specific enolase promoter inhibitor domain to prepare transgenic mice. See also, Wirak et al., *Science*, 253: 323-325
10 (1991). Still other models have been produced by Intracranial administration of the β /A4 peptide directly to animals (Tate et al., *Bull. Clin. Neurosci.*, 56: 131-139 (1991)).

15 Without further elaboration, it is believed that one skilled in the art can, using the preceding description, utilize the present invention to its fullest extent. The following examples are merely illustrative and not limitative of the remainder of the disclosure in any way whatsoever.

EXAMPLES

As noted above, A4 is intended to be the same as β /A4, throughout the examples. The peptides used in the following Examples have the following structures:

5 **A4-O (peptides 1-28), SEQ ID NO:7:**

N-Asp-Ala-Glu-Phe-Arg-His-Asp-Ser-Gly-Tyr-Glu-Val-His-His-Gln-Lys-Leu-Val-Phe-Phe-Ala-Glu-Asp-Val-Gly-Ser-Ser-Ala-COOH

10 The A4-O(1-28) polypeptide that was reported in Masters, et al. *Proc. Nat'l. Acad. Sci. U.S.A.*, 82: 4245-4249 (1985) is the first 28 amino acids of the 4.2 Kd peptide derived from senile plaque cores of an AD brain. Masters, et al. have also shown that the naturally occurring peptide aggregates even in denaturing gels.

15 The A4-O(1-28) sequence of this invention was synthesized by Biosearch in San Rafael, CA. The underlined amino acids differ from A4-P(1-28), as shown below.

A4-H (peptides 1-28):

20 The A4-H peptide is the same as A4-O(1-28) except that it was synthesized by the Harvard Microchemistry Laboratory.

A4-P (peptides 1-28), SEQ ID NO:8:

N-Asp-Ala-Glu-Phe-Arg-His-Asp-Ser-Gly-Tyr-Gln-Val-His-His-Gln-Lys-Leu-Val-Phe-Phe-Ala-Glu-Asp-Val-Gly-Ser-Asn-Lys-COOH

25

30 This sequence was reported by Glenner and Wong, *supra*, (1984) and derived from vascular amyloid of the AD brain and from a Down Syndrome brain. Three of 28 amino acids are different from the A4-O/A4-H peptides (underlined).

A4-B (peptides 1-28), SEQ ID NO:9:

N-Asp-Ala-Glu-Phe-Arg-His-Asp-Ser-Gly-Tyr-Glu-Val-His-His-Gln-Lys-Leu-Val-Phe-Phe-Ala-Glu-Asp-Val-Gly-Ser-Asn-Lys-COOH

35 This sequence was obtained from Bachem and is the 28 amino acid structure that is commonly determined from

molecular cloning studies (Kang, et al., *Nature (London)* 325: 733-736 (1987)). Unlike the Glenner and Wong, *supra*, sequence (A4-P(1-28)), it has Glu, not Gln, at position 11. And, unlike A4-O/A4-H, it has Asn-Lys and not Ser-Ala at the C-terminus.

Op1 (peptides 1-10), SEQ ID NO:10:

N-Asp-Ala-Glu-Phe-Arg-His-Asp-Ser-Gly-Tyr-COOH

A4(1-10) consists of the first 10 amino acids of the amyloid peptide derived from any source and is described in U.S. Patent No. 4,666,829 by Glenner et al. Thus far, this sequence appears conserved in all reports on amyloid that is derived from non-Familial AD cases. The A4-(1-10) antigen used in the present studies was synthesized by the Harvard Microchemistry Laboratory.

Summary of sequence variations: dashed line indicates sequence conservation among the peptides shown.

A4-O (1-28):	N-----Glu-----Ser-Ala-	(Masters)
A4-H (1-28):	N-----Glu-----Ser-Ala-	(Masters)
A4-P (1-28):	N-----Gln-----Asn-Lys-	(Glenner)
A4-B (1-28):	N-----Glu-----Asn-Lys-	(Kang)
Op1	: N-----	

Example 1. Self-aggregation of the A4-O amyloid peptide in SDS/urea acrylamide gels

The synthetic β -amyloid 28-mer polypeptide, A4-O (Masters, et al., *supra.*) was analyzed by polyacrylamide gel electrophoresis (PAGE) procedures (Brown, et al., *J. Neurochem.*, 40: 299-308 (1983)) and was noted to have unusual aggregation properties. The peptide was dissolved in a PAGE sample buffer containing SDS and urea and was electrophoresed on a 10% gel containing SDS (See description of Figure 2). After staining with Coomassie blue, the peptide appeared as a broad band at approximately 23-25 kd and a narrow band that migrated at the get front during electrophoresis (See Figure 2). The higher molecular weight species appeared to be an aggregate since it was eliminated by adding urea to the separating get and, subsequently, a 3-4 kd band was obtained (not shown). Polyclonal antiserum to the 28-mer was prepared and applied to nitrocellulose blots of an overloaded gel. The latter contained a series of aggregated peptides of various apparent molecular weights, all of which reacted with the antiserum. Thus, the synthetic 28-mer had aggregational properties not unlike the naturally occurring A4-O amyloid protein of 4 kd (Masters, et al., *supra.*).

Applicants' studies demonstrating the aggregation properties of the A4-O peptide were previously reported (Salim, et al., "Molecular Cloning of Amyloid cDNA from Alzheimer Brain Messenger RNA" in *Familial Alzheimer's Disease*, J.P. Blass et al. eds., Marcel Dekker, NY pp 153-165 (1988)).

Based upon the results shown in Figure 2, applicants concluded that even in the presence of strong denaturing agents and after electrophoresis, A4-O strongly bound to itself.

Example 2: Self-aggregation of A4-O peptide on highly cross-linked SDS/urea acrylamide gels

Applicants obtained confirmatory data using the highly cross-linked acrylamide gel system described by
5 Honda and Marotta, *Neurochem. Res.*, 17: 367-374 (1992).

When analyzed by this gel system containing SDS and urea, the synthetic peptide A4-O migrated as a broad series of bands below an apparent molecular weight of 15kDa (data not shown). However, when 6M urea was added
10 to the PAGE system the peptide appeared as a sharp single band of 15kDa (Figure 3) and smaller size bands were not observed even after silver staining (data not shown). By contrast, peptide P2(413-429), used as a control and corresponding to an extracytoplasmic region of the B/A4
15 precursor protein, migrated with the bromphenol blue dye front on both SDS-PAGE and SDS/urea-PAGE systems (Figure 3, lane 4). Since the theoretical molecular weight of the 28 amino acid peptide A4-O is 3,178 Da the results indicate that the band of 15kDa is an aggregate.
20 Migration of A4-O peptide bands on both gel systems was not affected by 2-ME nor by pre-treatment with 80% formic acid (data not shown).

The 15kDa was visible after peptide A4-O were treated for 5 minutes at 95°C prior to electrophoresis (Figure 3,
25 lane 1). When boiling time was increased to 30 minutes or 60 minutes the aggregate partly dissociated to a smaller size (Figure 3, lanes 2 and 3). This dissociation was not dependent on the presence of SDS and 2-ME in the sample buffer but rather on the time of heat
30 denaturation. These data were previously reported (Honda and Marotta, *supra*).

Applicants concluded that Figure 3 confirms that even in the presence of strong denaturing agents and heat treatment after electrophoresis, A4-O strongly bound to
35 itself.

Example 3: Self-aggregation of A4 peptides on immunoblots

Due to the desirability of obtaining a quantitative assay for the selection of B-amyloid polypeptides for the composition and methods of the present invention, applicants elected to use quantitative slot blots to test aggregation of peptides rather than tissue slices. The general immunoblotting procedure utilizing A4 peptides attached to a solid support and detectable by applied anti-amyloid antibodies was reported earlier (Majocha, et al., *supra*, (1988). In all cases, the monoclonal antibody used to detect A4 aggregates was 10H3 (2 ug/ml).

One microgram of each of the indicated peptides were added overnight at room temperature to Millipore P filter paper to which A4-O was attached. The peptides were dissolved in ICC buffer: 2% BSA, 0.3M NaCl, 20mM Tris, 0.01% Triton. The blots were immunoprocessed (Majocha, et al. *supra*.) and then optically scanned for density; the areas under the curves were integrated by means of an LKB Laser Densitometer.

Peptides A4-O, A4-H and Op1 were applied to filters to which was bound peptide A4-O, the antigen used to prepare mab 10H3. The experiment was designed to test the competence of each of the applied peptides to bind to the bound peptide. While A4-O and A4-H have the same primary structure, it has been noted that peptides with identical sequences that are obtained from different sources may have non-identical properties. (See Figure 4).

The density of staining (the optical density of the immunoreaction product) is quantitated in Figures 5, 6 and 7. The OD is a measure of the extent of the aggregation since it will be related to the antibody concentration and thus the color reaction.

The density values shown in Figure 5 were obtained by densitometric scanning of the reaction product on

blots from which the control value (no primary antibody) was subtracted. A further control was one in which the mab 10H3 was added to blots containing Op1 in the absence of added exogenous peptide. This control value
5 represents the antibody-antigen reaction without interference from added peptides.

Based upon the results presented in Figure 5, applicants concluded that A4-O bound to itself with at an optimal concentration of 5.0 ug/ml.

10 Example 4: Self-aggregation of A4-H peptides on immunoblots

The experiment of Example 3 was repeated except that the exogenous peptide was A4-H. The data are shown in Figure 6 and based upon these results, applicants
15 concluded that A4-H bound to A4-O at an optimal concentration of 2.5 ug/ml.

Example 5: Self-aggregation of Op1 peptides on immunoblots

The experiment of Example 3 was repeated except that the exogenous peptide was Op1. The data are shown in Figure 7 and based upon these results, applicants concluded that Op1 bound to A4-O at an optimal concentration of 2.5 ug/ml.

Based upon the results presented in Figures 5, 6 and 7, applicants concluded that three peptides bound the filter-bound A4-O peptide and increased the extent to which 10H3 reacted. The reaction is concentration-dependent. The three peptides, A4-O, A4-H and Op1, aggregated to the attached A4-O. The Op1 10-mer reacted nearly as well or better, at 2.5 ug, as the larger 28-mers.

Example 6: Specificity of 10H3 for both A4-O and Op1

The results shown in Figure 7 indicate that a small peptide, a ten-mer, was able to bind at least as well as 1-28-mers to an A4 substrate.

Thus, this assay, which measures the optical density of the reaction product between the added 10H3 mab and the Op1 peptide on the solid surface, reflected the presence of the exogenous peptide, as applicants previously demonstrated for the reaction between 10H3 and A4-O.

With respect to Op1, additional studies were carried out to confirm the reactivity of 10H3. On separate solid supports (Millipore P paper) either the A4-O antigen (2ug/slot) or the Op1 antigen (2 ug/slot) were absorbed using a slot blot apparatus. The results are shown in Figure 8, as follows:

Reactivity of 10H3 towards A4-O (upper panel):

Blot no:

1. Immunostain lacking the primary antibody (10H3) showed no reactivity with the blot, as expected.

-30-

2. 10H3 was very strongly reactive with its own antigen, A4-O.
 3. Soluble A4-O antigen added to the mix caused inhibition of 10H3 towards A4-O.
 - 5 4. Soluble Op1 added to the mix caused inhibition of 10H3 towards A4-O.
 5. 10H3 was reactive towards the Op1 antigen.
 6. Soluble Op1 added to the mix showed inhibition of 10H3 towards Op1.
- 10 The slot blots were quantified by densitometry and numerical values were obtained that indicated the extent of the reaction between 10H3 and antigens. These values are given below in Table I in which each numbered item refers to the blot number in Figure 8 and the description
- 15 given above:

Table I: Optical Density of Reaction Between 10H3 and Either A4-O or Op1 Antigens in Blots of Figure 7

Slot Number:	1	2	3	4	5	6
OD Units:	0.03	0.70	0.31	0.18	0.16	0.07

Based upon the results presented in Figure 8 and Table I, applicants concluded that 10H3 is reactive with its own A4-O antigen as well as with the Op1 peptide.

-31-

SEQUENCE LISTING

(1) GENERAL INFORMATION:

- (i) APPLICANT: THE MIRIAM HOSPITAL
- (ii) TITLE OF INVENTION: Composition and Method for in Vivo Imaging of Amyloid Deposits
- (iii) NUMBER OF SEQUENCES: 13
- (iv) CORRESPONDENCE ADDRESS:
 - (A) ADDRESSEE: Foley & Lardner
 - (B) STREET: 3000 K Street, N.W., Suite 500
 - (C) CITY: Washington, D.C.
 - (E) COUNTRY: USA
 - (F) ZIP: 20007-5109
- (v) COMPUTER READABLE FORM:
 - (A) MEDIUM TYPE: Floppy disk
 - (B) COMPUTER: IBM PC compatible
 - (C) OPERATING SYSTEM: PC-DOS/MS-DOS
 - (D) SOFTWARE: PatentIn Release #1.0, Version #1.25
- (vi) CURRENT APPLICATION DATA:
 - (A) APPLICATION NUMBER:
 - (B) FILING DATE: 27 May 1994
 - (C) CLASSIFICATION:
- (viii) ATTORNEY/AGENT INFORMATION:
 - (A) NAME: SAXE, Bernhard D.
 - (B) REGISTRATION NUMBER: 28,665
 - (C) REFERENCE/DOCKET NUMBER: 57548/103/MIHO
- (ix) TELECOMMUNICATION INFORMATION:
 - (A) TELEPHONE: (202)672-5300
 - (B) TELEFAX: (202)672-5399
 - (C) TELEX: 904136

(2) INFORMATION FOR SEQ ID NO:1:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 28 amino acids
 - (B) TYPE: amino acid
 - (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

Asp	Ala	Glu	Phe	Arg	His	Asp	Ser	Gly	Tyr	Gln	Val	His	His	Gln	Lys
1				5					10					15	
Leu	Val	Phe	Phe	Ala	Glu	Asp	Val	Gly	Ser	Asn	Lys				
			20					25							

(2) INFORMATION FOR SEQ ID NO:2:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 43 amino acids
 - (B) TYPE: amino acid
 - (D) TOPOLOGY: linear

-32-

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

Asp Ala Glu Phe Arg His Asp Ser Gly Tyr Glu Val His His Gln Lys
 1 5 10 15
 Leu Val Phe Phe Ala Glu Asp Val Gly Ser Asn Lys Gly Ala Ile Ile
 20 25 30
 Gly Leu Met Val Gly Gly Val Val Ile Ala Thr
 35 40

(2) INFORMATION FOR SEQ ID NO:3:

- (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 52 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

Asp Ala Glu Phe Arg His Asp Ser Gly Tyr Glu Val His His Gln Lys
 1 5 10 15
 Leu Val Phe Phe Ala Glu Asp Val Gly Ser Asn Lys Gly Ala Ile Ile
 20 25 30
 Gly Leu Met Val Gly Gly Val Val Ile Ala Thr Val Ile Val Ile Thr
 35 40 45
 Leu Val Met Leu
 50

(2) INFORMATION FOR SEQ ID NO:4:

- (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 99 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:

Asp Ala Glu Phe Arg His Asp Ser Gly Tyr Glu Val His His Gln Lys
 1 5 10 15
 Leu Val Phe Phe Ala Glu Asp Val Gly Ser Asn Lys Gly Ala Ile Ile
 20 25 30
 Gly Leu Met Val Gly Gly Val Val Ile Ala Thr Val Ile Val Ile Thr
 35 40 45
 Leu Val Met Leu Lys Lys Lys Gln Tyr Thr Ser Ile His His Gly Val
 50 55 60
 Val Glu Val Asp Ala Ala Val Thr Pro Glu Glu Arg His Leu Ser Lys
 65 70 75 80
 Met Gln Gln Asn Gly Tyr Glu Asn Pro Thr Tyr Lys Phe Phe Glu Gln
 85 90 95
 Met Gln Asn

(2) INFORMATION FOR SEQ ID NO:5:

-33-

- (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 53 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:5:

```

Lys Thr Glu Glu Ile Ser Glu Val Lys Met Asp Ala Glu Phe Arg His
1           5           10           15
Asp Ser Gly Tyr Glu Val His His Gln Lys Leu Val Phe Phe Ala Glu
20           25           30
Asp Val Gly Ser Asn Lys Gly Ala Ile Ile Gly Leu Met Val Gly Gly
35           40           45
Val Val Ile Ala Thr
50

```

(2) INFORMATION FOR SEQ ID NO:6:

- (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 45 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

(ix) FEATURE:

- (A) NAME/KEY: Modified-site
 (B) LOCATION: 1
 (D) OTHER INFORMATION: /note= "Xaa at position 1
 corresponds to 1 or more APP amino acids which are
 not adjacent to B/A4 in nature."

(ix) FEATURE:

- (A) NAME/KEY: Modified-site
 (B) LOCATION: 45
 (D) OTHER INFORMATION: /note= "Xaa at position 45
 corresponds to 1 or more APP amino acids which are
 not adjacent to B/A4 in nature."

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:6:

```

Xaa Asp Ala Glu Phe Arg His Asp Ser Gly Tyr Glu Val His His Gln
1           5           10           15
Lys Leu Val Phe Phe Ala Glu Asp Val Gly Ser Asn Lys Gly Ala Ile
20           25           30
Ile Gly Leu Met Val Gly Gly Val Val Ile Ala Thr Xaa
35           40           45

```

(2) INFORMATION FOR SEQ ID NO:7:

- (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 28 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

- (vii) IMMEDIATE SOURCE:
 (B) CLONE: A4-O

-34-

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:7:

Asp Ala Glu Phe Arg His Asp Ser Gly Tyr Glu Val His His Gln Lys
1 5 10 15
Leu Val Phe Phe Ala Glu Asp Val Gly Ser Ser Ala
20 25

(2) INFORMATION FOR SEQ ID NO:8:

- (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 28 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

- (vii) IMMEDIATE SOURCE:
(B) CLONE: A4-P

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:8:

Asp Ala Glu Phe Arg His Asp Ser Gly Tyr Gln Val His His Gln Lys
1 5 10 15
Leu Val Phe Phe Ala Glu Asp Val Gly Ser Asn Lys
20 25

(2) INFORMATION FOR SEQ ID NO:9:

- (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 28 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

- (vii) IMMEDIATE SOURCE:
(B) CLONE: A4-B

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:9:

Asp Ala Glu Phe Arg His Asp Ser Gly Tyr Glu Val His His Gln Lys
1 5 10 15
Leu Val Phe Phe Ala Glu Asp Val Gly Ser Asn Lys
20 25

(2) INFORMATION FOR SEQ ID NO:10:

- (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 10 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

- (vii) IMMEDIATE SOURCE:
(B) CLONE: Op1

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:10:

Asp Ala Glu Phe Arg His Asp Ser Gly Tyr
1 5 10

(2) INFORMATION FOR SEQ ID NO:11:

-35-

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 3353 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: double
- (D) TOPOLOGY: linear

(ix) FEATURE:

- (A) NAME/KEY: CDS
- (B) LOCATION: 147..2234

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:11:

```

AGTTTCCTCG GCAGCGGTAG GCGAGAGCAC GCGGAGGAGC GTGCGCGGGG CCCCAGGAGA      60
CGGCGGCGGT GCGGCGCGG GCAGAGCAAG GACGCGGCGG ATCCCACTCG CACAGCAGCG      120
CACTCGGTGC CCCGCGCAGG GTCGCG ATG CTG CCC GGT TTG GCA CTG CTC CTG      173
               Met Leu Pro Gly Leu Ala Leu Leu Leu
                1                               5

CTG GCC GCC TGG ACG GCT CGG GCG CTG GAG GTA CCC ACT GAT GGT AAT      221
Leu Ala Ala Trp Thr Ala Arg Ala Leu Glu Val Pro Thr Asp Gly Asn
 10                      15                      20                      25

GCT GGC CTG CTG GCT GAA CCC CAG ATT GCC ATG TTC TGT GGC AGA CTG      269
Ala Gly Leu Leu Ala Glu Pro Gln Ile Ala Met Phe Cys Gly Arg Leu
                30                      35                      40

AAC ATG CAC ATG AAT GTC CAG AAT GGG AAG TGG GAT TCA GAT CCA TCA      317
Asn Met His Met Asn Val Gln Asn Gly Lys Trp Asp Ser Asp Pro Ser
                45                      50                      55

GGG ACC AAA ACC TGC ATT GAT ACC AAG GAA GGC ATC CTG CAG TAT TGC      365
Gly Thr Lys Thr Cys Ile Asp Thr Lys Glu Gly Ile Leu Gln Tyr Cys
                60                      65                      70

CAA GAA GTC TAC CCT GAA CTG CAG ATC ACC AAT GTG GTA GAA GCC AAC      413
Gln Glu Val Tyr Pro Glu Leu Gln Ile Thr Asn Val Val Glu Ala Asn
                75                      80                      85

CAA CCA GTG ACC ATC CAG AAC TGG TGC AAG CGG GGC CGC AAG CAG TGC      461
Gln Pro Val Thr Ile Gln Asn Trp Cys Lys Arg Gly Arg Lys Gln Cys
                90                      95                      100                      105

AAG ACC CAT CCC CAC TTT GTG ATT CCC TAC CGC TGC TTA GTT GGT GAG      509
Lys Thr His Pro His Phe Val Ile Pro Tyr Arg Cys Leu Val Gly Glu
                110                      115                      120

TTT GTA AGT GAT GCC CTT CTC GTT CCT GAC AAG TGC AAA TTC TTA CAC      557
Phe Val Ser Asp Ala Leu Leu Val Pro Asp Lys Cys Lys Phe Leu His
                125                      130                      135

CAG GAG AGG ATG GAT GTT TGC GAA ACT CAT CTT CAC TGG CAC ACC GTC      605
Gln Glu Arg Met Asp Val Cys Glu Thr His Leu His Trp His Thr Val
                140                      145                      150

GCC AAA GAG ACA TGC AGT GAG AAG AGT ACC AAC TTG CAT GAC TAC GGC      653
Ala Lys Glu Thr Cys Ser Glu Lys Ser Thr Asn Leu His Asp Tyr Gly
                155                      160                      165

ATG TTG CTG CCC TGC GGA ATT GAC AAG TTC CGA GGG GTA GAG TTT GTG      701
Met Leu Leu Pro Cys Gly Ile Asp Lys Phe Arg Gly Val Glu Phe Val
                170                      175                      180                      185

TGT TGC CCA CTG GCT GAA GAA AGT GAC AAT GTG GAT TCT GCT GAT GCG      749

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-36-

Cys	Cys	Pro	Leu	Ala	Glu	Glu	Ser	Asp	Asn	Val	Asp	Ser	Ala	Asp	Ala	
				190					195					200		
GAG	GAG	GAT	GAC	TCG	GAT	GTC	TGG	TGG	GGC	GGA	GCA	GAC	ACA	GAC	TAT	797
Glu	Glu	Asp	Asp	Ser	Asp	Val	Trp	Trp	Gly	Gly	Ala	Asp	Thr	Asp	Tyr	
			205					210					215			
GCA	GAT	GGG	AGT	GAA	GAC	AAA	GTA	GTA	GAA	GTA	GCA	GAG	GAG	GAA	GAA	845
Ala	Asp	Gly	Ser	Glu	Asp	Lys	Val	Val	Glu	Val	Ala	Glu	Glu	Glu	Glu	
		220					225					230				
GTG	GCT	GAG	GTG	GAA	GAA	GAA	GAA	GCC	GAT	GAT	GAC	GAG	GAC	GAT	GAG	893
Val	Ala	Glu	Val	Glu	Glu	Glu	Glu	Ala	Asp	Asp	Asp	Glu	Asp	Asp	Glu	
	235					240					245					
GAT	GGT	GAT	GAG	GTA	GAG	GAA	GAG	GCT	GAG	GAA	CCC	TAC	GAA	GAA	GCC	941
Asp	Gly	Asp	Glu	Val	Glu	Glu	Glu	Ala	Glu	Glu	Pro	Tyr	Glu	Glu	Ala	
	250				255					260					265	
ACA	GAG	AGA	ACC	ACC	AGC	ATT	GCC	ACC	ACC	ACC	ACC	ACC	ACC	ACA	GAG	989
Thr	Glu	Arg	Thr	Thr	Ser	Ile	Ala	Thr	Thr	Thr	Thr	Thr	Thr	Thr	Glu	
				270				275						280		
TCT	GTG	GAA	GAG	GTG	GTT	CGA	GTT	CCT	ACA	ACA	GCA	GCC	AGT	ACC	CCT	1037
Ser	Val	Glu	Glu	Val	Val	Arg	Val	Pro	Thr	Thr	Ala	Ala	Ser	Thr	Pro	
			285					290					295			
GAT	GCC	GTT	GAC	AAG	TAT	CTC	GAG	ACA	CCT	GGG	GAT	GAG	AAT	GAA	CAT	1085
Asp	Ala	Val	Asp	Lys	Tyr	Leu	Glu	Thr	Pro	Gly	Asp	Glu	Asn	Glu	His	
		300					305					310				
GCC	CAT	TTC	CAG	AAA	GCC	AAA	GAG	AGG	CTT	GAG	GCC	AAG	CAC	CGA	GAG	1133
Ala	His	Phe	Gln	Lys	Ala	Lys	Glu	Arg	Leu	Glu	Ala	Lys	His	Arg	Glu	
	315					320				325						
AGA	ATG	TCC	CAG	GTC	ATG	AGA	GAA	TGG	GAA	GAG	GCA	GAA	CGT	CAA	GCA	1181
Arg	Met	Ser	Gln	Val	Met	Arg	Glu	Trp	Glu	Glu	Ala	Glu	Arg	Gln	Ala	
	330				335				340					345		
AAG	AAC	TTG	CCT	AAA	GCT	GAT	AAG	AAG	GCA	GTT	ATC	CAG	CAT	TTC	CAG	1229
Lys	Asn	Leu	Pro	Lys	Ala	Asp	Lys	Lys	Ala	Val	Ile	Gln	His	Phe	Gln	
				350				355						360		
GAG	AAA	GTG	GAA	TCT	TTG	GAA	CAG	GAA	GCA	GCC	AAC	GAG	AGA	CAG	CAG	1277
Glu	Lys	Val	Glu	Ser	Leu	Glu	Gln	Glu	Ala	Ala	Asn	Glu	Arg	Gln	Gln	
			365				370						375			
CTG	GTG	GAG	ACA	CAC	ATG	GCC	AGA	GTG	GAA	GCC	ATG	CTC	AAT	GAC	CGC	1325
Leu	Val	Glu	Thr	His	Met	Ala	Arg	Val	Glu	Ala	Met	Leu	Asn	Asp	Arg	
		380				385						390				
CGC	CGC	CTG	GCC	CTG	GAG	AAC	TAC	ATC	ACC	GCT	CTG	CAG	GCT	GTT	CCT	1373
Arg	Arg	Leu	Ala	Leu	Glu	Asn	Tyr	Ile	Thr	Ala	Leu	Gln	Ala	Val	Pro	
	395					400					405					
CCT	CGG	CCT	CGT	CAC	GTG	TTC	AAT	ATG	CTA	AAG	AAG	TAT	GTC	CGC	GCA	1421
Pro	Arg	Pro	Arg	His	Val	Phe	Asn	Met	Leu	Lys	Lys	Tyr	Val	Arg	Ala	
	410				415					420				425		
GAA	CAG	AAG	GAC	AGA	CAG	CAC	ACC	CTA	AAG	CAT	TTC	GAG	CAT	GTG	CGC	1469
Glu	Gln	Lys	Asp	Arg	Gln	His	Thr	Leu	Lys	His	Phe	Glu	His	Val	Arg	
				430				435						440		
ATG	GTG	GAT	CCC	AAG	AAA	GCC	GCT	CAG	ATC	CGG	TCC	CAG	GTT	ATG	ACA	1517
Met	Val	Asp	Pro	Lys	Lys	Ala	Ala	Gln	Ile	Arg	Ser	Gln	Val	Met	Thr	
			445					450					455			

-37-

CAC CTC CGT GTG ATT TAT GAG CGC ATG AAT CAG TCT CTC TCC CTG CTC	1565
His Leu Arg Val Ile Tyr Glu Arg Met Asn Gln Ser Leu Ser Leu Leu	
460 465 470	
TAC AAC GTG CCT GCA GTG GCC GAG GAG ATT CAG GAT GAA GTT GAT GAG	1613
Tyr Asn Val Pro Ala Val Ala Glu Glu Ile Gln Asp Glu Val Asp Glu	
475 480 485	
CTG CTT CAG AAA GAG CAA AAC TAT TCA GAT GAC GTC TTG GCC AAC ATG	1661
Leu Leu Gln Lys Glu Gln Asn Tyr Ser Asp Asp Val Leu Ala Asn Met	
490 495 500 505	
ATT AGT GAA CCA AGG ATC AGT TAC GGA AAC GAT GCT CTC ATG CCA TCT	1709
Ile Ser Glu Pro Arg Ile Ser Tyr Gly Asn Asp Ala Leu Met Pro Ser	
510 515 520	
TTG ACC GAA ACG AAA ACC ACC GTG GAG CTC CTT CCC GTG AAT GGA GAG	1757
Leu Thr Glu Thr Lys Thr Thr Val Glu Leu Leu Pro Val Asn Gly Glu	
525 530 535	
TTC AGC CTG GAC GAT CTC CAG CCG TGG CAT TCT TTT GGG GCT GAC TCT	1805
Phe Ser Leu Asp Asp Leu Gln Pro Trp His Ser Phe Gly Ala Asp Ser	
540 545 550	
GTG CCA GCC AAC ACA GAA AAC GAA GTT GAG CCT GTT GAT GCC CGC CCT	1853
Val Pro Ala Asn Thr Glu Asn Glu Val Glu Pro Val Asp Ala Arg Pro	
555 560 565	
GCT GCC GAC CGA GGA CTG ACC ACT CGA CCA GGT TCT GGG TTG ACA AAT	1901
Ala Ala Asp Arg Gly Leu Thr Thr Arg Pro Gly Ser Gly Leu Thr Asn	
570 575 580 585	
ATC AAG ACG GAG GAG ATC TCT GAA GTG AAG ATG GAT GCA GAA TTC CGA	1949
Ile Lys Thr Glu Glu Ile Ser Glu Val Lys Met Asp Ala Glu Phe Arg	
590 595 600	
CAT GAC TCA GGA TAT GAA GTT CAT CAT CAA AAA TTG GTG TTC TTT GCA	1997
His Asp Ser Gly Tyr Glu Val His His Gln Lys Leu Val Phe Phe Ala	
605 610 615	
GAA GAT GTG GGT TCA AAC AAA GGT GCA ATC ATT GGA CTC ATG GTG GGC	2045
Glu Asp Val Gly Ser Asn Lys Gly Ala Ile Ile Gly Leu Met Val Gly	
620 625 630	
GGT GTT GTC ATA GCG ACA GTG ATC GTC ATC ACC TTG GTG ATG CTG AAG	2093
Gly Val Val Ile Ala Thr Val Ile Val Ile Thr Leu Val Met Leu Lys	
635 640 645	
AAG AAA CAG TAC ACA TCC ATT CAT CAT GGT GTG GTG GAG GTT GAC GCC	2141
Lys Lys Gln Tyr Thr Ser Ile His His Gly Val Val Glu Val Asp Ala	
650 655 660 665	
GCT GTC ACC CCA GAG GAG CGC CAC CTG TCC AAG ATG CAG CAG AAC GGC	2189
Ala Val Thr Pro Glu Glu Arg His Leu Ser Lys Met Gln Gln Asn Gly	
670 675 680	
TAC GAA AAT CCA ACC TAC AAG TTC TTT GAG CAG ATG CAG AAC TAGACCCCG	2241
Tyr Glu Asn Pro Thr Tyr Lys Phe Phe Glu Gln Met Gln Asn	
685 690 695	
CCACAGCAGC CTCTGAAGTT GGACAGCAAA ACCATTGCTT CACTACCCAT CGGTGTCCAT	2301
TTATAGAATA ATGTGGGAAG AAACAAACCC GTTTTATGAT TTACTCATTA TCGCCTTTTG	2361
ACAGCTGTGC TGTAACACAA GTAGATGCCT GAACCTGAAT TAATCCACAC ATCAGTAATG	2421
TATTCTATCT CTCTTTACAT TTTGGTCTCT ATACTACATT ATTAATGGGT TTTGTGTACT	2481

-38-

GTAAAGAATT TAGCTGTATC AAAC TAGTGC ATGAATAGAT TCTCTCCTGA TTATTTATCA 2541
CATAGCCCCCT TAGCCAGTTG TATATTATTC TTGTGGTTTG TGACCCAATT AAGTCCTACT 2601
TTACATATGC TTTAAGAATC GATGGGGGAT GCTTCATGTG AACGTGGGAG TTCAGCTGCT 2661
TCTCTGCGCT AAGTATTCCT TTCCTGATCA CTATGCATTT TAAAGTTAAA CATTTTAAAG 2721
TATTTTCAGAT GCTTTAGAGA GATTTTTTTT CCATGACTGC ATTTTACTGT ACAGATTGCT 2781
GCTTCTGCTA TATTTGTGAT ATAGGAATTA ACAGGATACA CACGTTTGTT TCTTCGTGCC 2841
TGTTTTATGT GCACACATTA GGCATTGAGA CTTCAAGCTT TTCTTTTTTT GTCCACGTAT 2901
CTTTGGGTCT TTGATAAAGA AAAGAATCCC TGTTTCATTGT AAGCACTTTT ACGGGGCGGG 2961
TGGGGAGGGG TGCTCTGCTG GTCTTCAATT ACCAAGAATT CTCCAAAACA ATTTTCTGCA 3021
GGATGATTGT ACAGAATCAT TGCTTATGAC ATGATCGCTT TCTACACTGT ATTACATAAA 3081
TAAATTAAAT AAAATAACCC CGGGCAAGAC TTTTCTTTGA AGGATGACTA CAGACATTAA 3141
ATAATCGAAG TAATTTTGGG TGGGGAGAAG AGGCAGATTC AATTTTCTTT AACCAGTCTG 3201
AAGTTTCATT TATGATACAA AAGAAGATGA AAATGGAAGT GGCAATATAA GGGGATGAGG 3261
AAGGCATGCC TGGACAAACC CTTCTTTTAA GATGTGTCTT CAATTTGTAT AAAATGGTGT 3321
TTTCATGTAA ATAAATACAT TCTTGAGGA GC 3353

(2) INFORMATION FOR SEQ ID NO:12:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 695 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:12:

Met Leu Pro Gly Leu Ala Leu Leu Leu Leu Ala Ala Trp Thr Ala Arg
1 5 10 15
Ala Leu Glu Val Pro Thr Asp Gly Asn Ala Gly Leu Leu Ala Glu Pro
20 25 30
Gln Ile Ala Met Phe Cys Gly Arg Leu Asn Met His Met Asn Val Gln
35 40 45
Asn Gly Lys Trp Asp Ser Asp Pro Ser Gly Thr Lys Thr Cys Ile Asp
50 55 60
Thr Lys Glu Gly Ile Leu Gln Tyr Cys Gln Glu Val Tyr Pro Glu Leu
65 70 75 80
Gln Ile Thr Asn Val Val Glu Ala Asn Gln Pro Val Thr Ile Gln Asn
85 90 95
Trp Cys Lys Arg Gly Arg Lys Gln Cys Lys Thr His Pro His Phe Val
100 105 110
Ile Pro Tyr Arg Cys Leu Val Gly Glu Phe Val Ser Asp Ala Leu Leu
115 120 125
Val Pro Asp Lys Cys Lys Phe Leu His Gln Glu Arg Met Asp Val Cys

-39-

130	135	140
Glu Thr His Leu His Trp 145 150	His Thr Val Ala Lys 155	Glu Thr Cys Ser Glu 160
Lys Ser Thr Asn Leu His Asp 165	Tyr Gly Met Leu Leu Pro 170	Cys Gly Ile 175
Asp Lys Phe Arg Gly Val Glu 180	Phe Val Cys Cys Pro Leu 185 190	Ala Glu Glu
Ser Asp Asn Val Asp Ser Ala 195 200	Asp Ala Glu Glu Asp 205	Ser Asp Val
Trp Trp Gly Gly Ala Asp 210 215	Thr Asp Tyr Ala Asp 220	Gly Ser Glu Asp Lys
Val Val Glu Val Ala Glu 225 230	Glu Glu Glu Val Ala Glu 235	Val Glu Glu Glu
Glu Ala Asp Asp Asp Glu 245	Asp Asp Glu Asp Gly 250	Val Glu Glu
Glu Ala Glu Glu Pro Tyr 260	Glu Glu Ala Thr Glu Arg 265 270	Thr Thr Ser Ile
Ala Thr Thr Thr Thr Thr 275 280	Thr Glu Ser Val Glu 285	Val Val Arg
Val Pro Thr Thr Ala Ala 290 295	Ser Thr Pro Asp Ala 300	Val Asp Lys Tyr Leu
Glu Thr Pro Gly Asp Glu 305 310	Asn Glu His Ala His 315	Phe Gln Lys Ala Lys
Glu Arg Leu Glu Ala Lys 325	His Arg Glu Arg Met 330	Ser Gln Val Met Arg
Glu Trp Glu Glu Ala Glu 340	Arg Gln Ala Lys Asn 345	Leu Pro Lys Ala Asp
Lys Lys Ala Val Ile Gln 355	His Phe Gln Glu Lys 360 365	Val Glu Ser Leu Glu
Gln Glu Ala Ala Asn Glu 370 375	Arg Gln Gln Leu Val 380	Glu Thr His Met Ala
Arg Val Glu Ala Met Leu 385 390	Asn Asp Arg Arg Arg 395	Leu Ala Leu Glu Asn
Tyr Ile Thr Ala Leu Gln 405	Ala Val Pro Pro Arg 410	Pro Arg His Val Phe
Asn Met Leu Lys Lys Tyr 420	Val Arg Ala Glu Gln 425	Lys Asp Arg Gln His
Thr Leu Lys His Phe Glu 435	His Val Arg Met Val 440	Asp Pro Lys Lys Ala
Ala Gln Ile Arg Ser Gln 450 455	Val Met Thr His Leu 460	Arg Val Ile Tyr Glu
Arg Met Asn Gln Ser Leu 465 470	Ser Leu Leu Tyr Asn 475	Val Pro Ala Val Ala
Glu Glu Ile Gln Asp Glu 485	Val Asp Glu Leu Leu 490	Gln Lys Glu Gln Asn
		495

-40-

Tyr Ser Asp Asp Val Leu Ala Asn Met Ile Ser Glu Pro Arg Ile Ser
 500 505 510
 Tyr Gly Asn Asp Ala Leu Met Pro Ser Leu Thr Glu Thr Lys Thr Thr
 515 520 525
 Val Glu Leu Leu Pro Val Asn Gly Glu Phe Ser Leu Asp Asp Leu Gln
 530 535 540
 Pro Trp His Ser Phe Gly Ala Asp Ser Val Pro Ala Asn Thr Glu Asn
 545 550 555 560
 Glu Val Glu Pro Val Asp Ala Arg Pro Ala Ala Asp Arg Gly Leu Thr
 565 570 575
 Thr Arg Pro Gly Ser Gly Leu Thr Asn Ile Lys Thr Glu Glu Ile Ser
 580 585 590
 Glu Val Lys Met Asp Ala Glu Phe Arg His Asp Ser Gly Tyr Glu Val
 595 600 605
 His His Gln Lys Leu Val Phe Phe Ala Glu Asp Val Gly Ser Asn Lys
 610 615 620
 Gly Ala Ile Ile Gly Leu Met Val Gly Gly Val Val Ile Ala Thr Val
 625 630 635 640
 Ile Val Ile Thr Leu Val Met Leu Lys Lys Lys Gln Tyr Thr Ser Ile
 645 650 655
 His His Gly Val Val Glu Val Asp Ala Ala Val Thr Pro Glu Glu Arg
 660 665 670
 His Leu Ser Lys Met Gln Gln Asn Gly Tyr Glu Asn Pro Thr Tyr Lys
 675 680 685
 Phe Phe Glu Gln Met Gln Asn
 690 695

(2) INFORMATION FOR SEQ ID NO:13:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 28 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

(ix) FEATURE:

- (A) NAME/KEY: Modified-site
- (B) LOCATION: 11
- (D) OTHER INFORMATION: /note= ""Xaa at Position 11 is either Glu or Gln.""

(ix) FEATURE:

- (A) NAME/KEY: Modified-site
- (B) LOCATION: 27
- (D) OTHER INFORMATION: /note= ""Xaa at position 27 is either Ser or Asn.""

(ix) FEATURE:

- (A) NAME/KEY: Modified-site
- (B) LOCATION: 28
- (D) OTHER INFORMATION: /note= ""Xaa at position 28 is either Ala or Lys.""

-41-

(x1) SEQUENCE DESCRIPTION: SEQ ID NO:13:

Asp	Ala	Glu	Phe	Arg	His	Asp	Ser	Gly	Tyr	Xaa	Val	His	His	Gln	Lys
1				5				10						15	

Leu	Val	Phe	Phe	Ala	Glu	Asp	Val	Gly	Ser	Xaa	Xaa
			20					25			

What Is Claimed Is:

1. An amyloid binding composition for *in vivo* imaging of amyloid deposits comprising:

- (a) a labeled amyloid protein or variant thereof that binds to amyloid deposits *in vivo*; and
- (b) a pharmaceutically acceptable carrier.

2. The composition of claim 1, wherein said amyloid protein is β -amyloid polypeptide or a variant thereof.

3. The composition of claim 2, wherein said β -amyloid polypeptide variant has the following amino acid sequence (SEQ ID NO:13):

N-Asp-Ala-Glu-Phe-Arg-His-Asp-Ser-Gly-Tyr-X-Val-His-His-Gln-Lys-Leu-Val-Phe-Phe-Ala-Glu-Asp-Val-Gly-Ser-Y1-Y2-COOH;

wherein X is either Glu or Gln; Y1 is either Ser or Asn; and Y2 is either Ala or Lys.

4. The composition of claim 3, wherein said β -amyloid polypeptide variant is selected from the group consisting of (1) a variant wherein when X is Glu, Y1 is Ser and Y2 is Ala, (2) a variant wherein when X is Glu, Y1 is Asn and Y2 is Lys, and (3) a variant wherein when X is Gln, Y1 is Asn and Y2 is Lys.

5. The composition of claim 2, wherein said β -amyloid polypeptide or variant thereof has an amino acid sequence selected from the following group of amino acid sequences:

(A) (SEQ ID NO:2) Asp-Ala-Glu-Phe-Arg-His-Asp-Ser-Gly-Tyr-Glu-Val-His-His-Gln-Lys-Leu-Val-Phe-Phe-Ala-Glu-Asp-Val-Gly-Ser-Asn-Lys-Gly-Ala-Ile-Ile-Gly-Leu-Met-Val-Gly-Gly-Val-Val-Ile-Ala-Thr;

(B) (SEQ ID NO:3) Asp-Ala-Glu-Phe-Arg-His-Asp-Ser-Gly-Tyr-Glu-Val-His-His-Gln-Lys-Leu-Val-Phe-Phe-Ala-Glu-Asp-Val-Gly-Ser-Asn-Lys-Gly-Ala-Ile-Ile-Gly-Leu-Met-Val-

-43-

- Gly-Gly-Val-Val-Ile-Ala-Thr-Val-Ile-Val-Ile-Thr-Leu-Val-Met-Leu;
- (C) (SEQ ID NO:4) Asp-Ala-Glu-Phe-Arg-His-Asp-Ser-Gly-Tyr-Glu-Val-His-His-Gln-Lys-Leu-Val-Phe-Phe-Ala-Glu-Asp-Val-Gly-Ser-Asn-Lys-Gly-Ala-Ile-Ile-Gly-Leu-Met-Val-Gly-Gly-Val-Val-Ile-Ala-Thr-Val-Ile-Val-Ile-Thr-Leu-Val-Met-Leu-Lys-Lys-Lys-Gln-Tyr-Thr-Ser-Ile-His-His-Gly-Val-Val-Glu-Val-Asp-Ala-Ala-Val-Thr-Pro-Glu-Glu-Arg-His-Leu-Ser-Lys-Met-Gln-Gln-Asn-Gly-Tyr-Glu-Asn-Pro-Thr-Tyr-Lys-Phe-Phe-Glu-Gln-Met-Gln-Asn;
- (D) (SEQ ID NO:5) Lys-Thr-Glu-Glu-Ile-Ser-Glu-Val-Lys-Met-Asp-Ala-Glu-Phe-Arg-His-Asp-Ser-Gly-Tyr-Glu-Val-His-His-Gln-Lys-Leu-Val-Phe-Phe-Ala-Glu-Asp-Val-Gly-Ser-Asn-Lys-Gly-Ala-Ile-Ile-Gly-Leu-Met-Val-Gly-Gly-Val-Val-Ile-Ala-Thr;
- (E) (SEQ ID NO:6) X-Asp-Ala-Glu-Phe-Arg-His-Asp-Ser-Gly-Tyr-Glu-Val-His-His-Gln-Lys-Leu-Val-Phe-Phe-Ala-Glu-Asp-Val-Gly-Ser-Asn-Lys-Gly-Ala-Ile-Ile-Gly-Leu-Met-Val-Gly-Gly-Val-Val-Ile-Ala-Thr-Y,

wherein X and Y are one or more APP amino acids which are not adjacent to β /A4 in the nature;

and

(F) any fragment of (A)-(E), wherein said fragment is large enough to bind amyloid deposit *in vivo*.

6. The composition of claim 1, wherein said labeled amyloid protein is radiolabeled amyloid protein.

7. The composition of claim 1, wherein said radiolabeled amyloid protein is Technetium 99m-labeled amyloid protein.

8. An *in vivo* method for detecting amyloid deposits in a subject comprising the steps of

(a) administering to a subject a detectable quantity of an amyloid binding composition comprising a labeled amyloid protein or variant thereof and a pharmaceutically acceptable carrier; and

(b) detecting the binding of the labeled protein or variant thereof to the amyloid deposit.

9. The method of claim 8, wherein said amyloid protein is the β -amyloid polypeptide or variant thereof.

10. The method of claim 8, wherein said amyloid protein is radiolabeled.

11. The method of claim 10, wherein said detecting involves radioactive imaging.

12. The method of claim 8, wherein said administering is selected from the group consisting of intravenous injection, intraventricular injection and a combination of both intravenous and intraventricular injection.

13. The method of claim 8, wherein said amyloid deposits are located in the brain of a subject.

14. A method of diagnosing an amyloidosis-associated disease by detecting amyloid deposits in a subject suspected of having amyloid deposits, said method comprising the steps of:

(a) administering to a subject a detectable quantity of an amyloid binding composition comprising a labeled amyloid protein or variant thereof and a pharmaceutically acceptable carrier; and

(b) detecting the binding of said labeled protein to said amyloid deposit.

15. The method of claim 14, wherein said amyloidosis-associated disease is selected from the group consisting of Alzheimer's Disease and Down Syndrome.

1/10

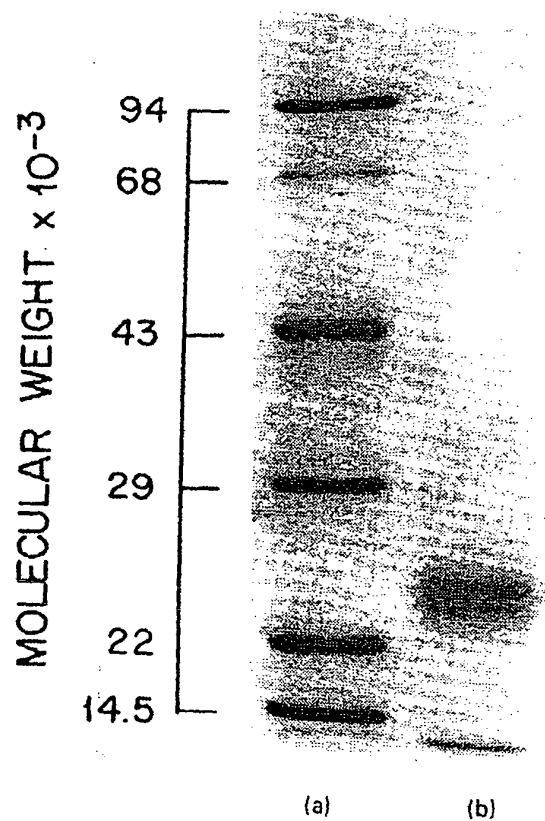
FIG. 1

Clinical Association	Notation	Protein Type
ACQUIRED SYSTEMIC AMYLOIDOSIS		
Immunoglobulin light-chain (primary), Multiple myeloma	AL	Light chain, type subtype
Reactive (secondary)	AA	Protein A
Hemodialysis amyloidosis	AH	β_2 microglobulin
HEREDOFAMILIAL		
Polyneuropathy	AF	Prealbumin, variant
Familial Mediterranean fever	AA	Protein A
ORGAN-LIMITED		
Hereditary Icelandic Congophilic angiopathy	AC _{vC}	Cystatin C, variant
Alzheimer's disease: vessels and plaques	AC _{vβ1} AC _{pβ}	β protein
Senile cardiac		
LOCALIZED ENDOCRINE		
Pancreatic islet	AE _f	Islet amyloid protein [IAP]
Medullary thyroid carcinoma	AE _t	Precalcitonin

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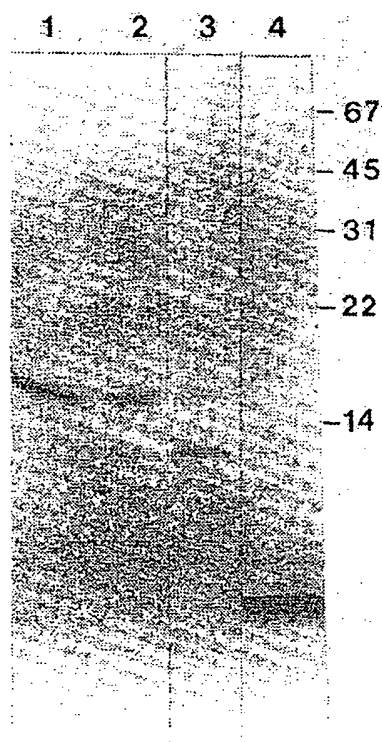
2/10

FIG. 2



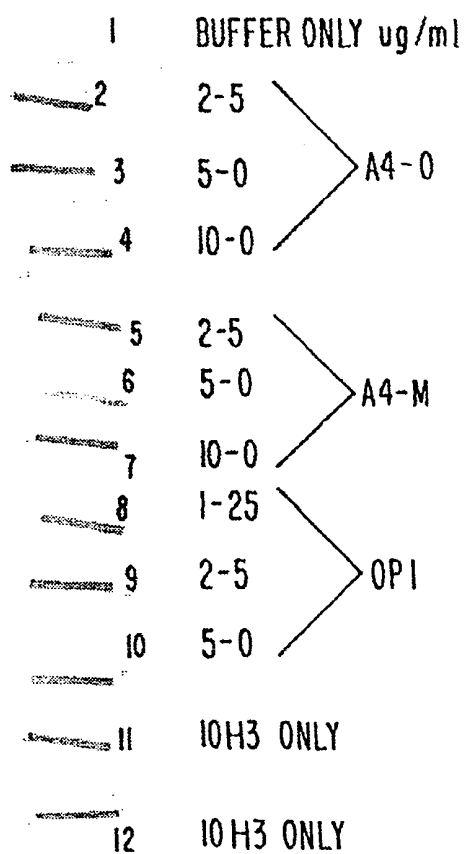
3/10

FIG. 3



4/10

FIG. 4

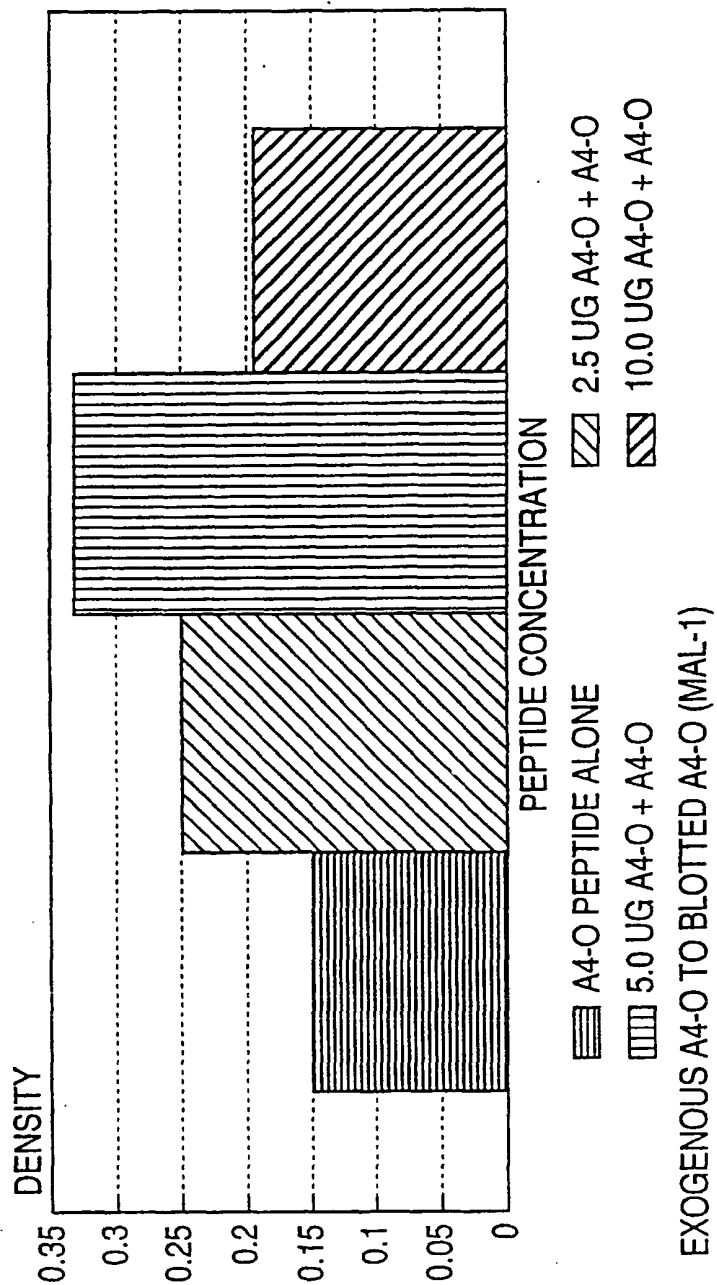


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5/10

FIG. 5

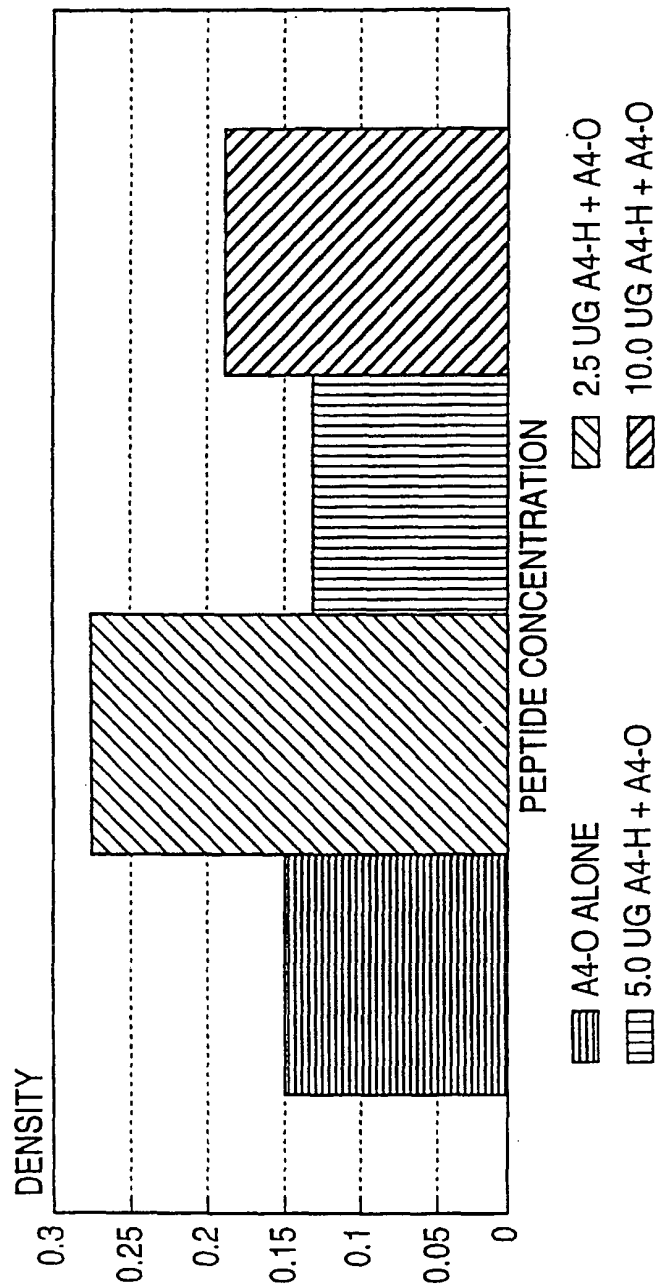
A4-O IMMUNOBLOT
AGGREGATION OF A4-O + A4-O



6/10

FIG. 6

A4-O IMMUNOBLOT
AGGREGATION OF A4-O + A4-H

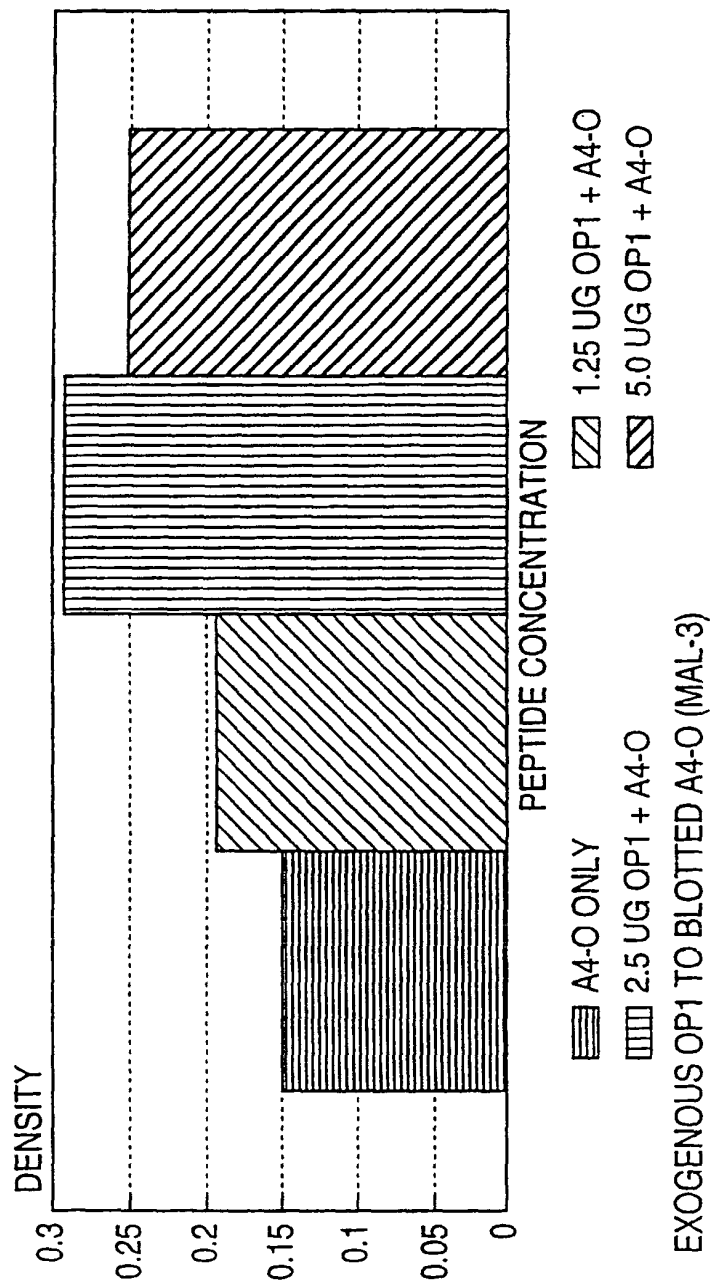


EXOGENOUS A4-H TO BLOTTED A4-O (MAL-2)

7/10

FIG. 7

A4-O IMMUNOBLOT
AGGREGATION OF A4-O + OP1



8/10

FIG. 8

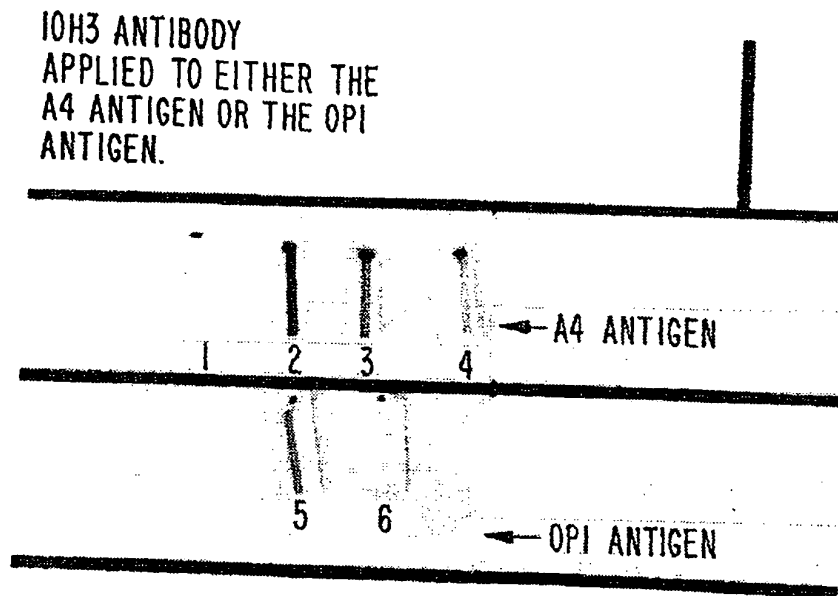


FIG. 9A

9/10

AGTTTCCTCGGCAGCGGTAGGCGAGA -121
2 GCACGCGGAGGAGCGTGC GCGGGGCCCCGGGAGACGGCGGCGGTGGCGGCGCGGGCAGAG -61
3 CAAGGACGCGGCGGATCCCACTCGCACAGCAGCGCACTCGGTGCCCCGCGCAGGGTCGCG -1
4 ATGCTGCCCCGTTTGGCACTGCTCCTGCTGGCCGCTGGACGGCTCGGGCGCTGGAGGTA 60
M L P G L A L L L A A W T A R A L E V
1 10 20
5 CCCACTGATGGTAATGCTGGCCTGCTGGCTGAACCCAGATTGCCATGTTCTGTGGCAGA 120
P T D G N A G L L A E P Q I A M F C G R
30 40
6 CTGAACATGCACATGAATGTCCAGAATGGGAAGTGGGATTTCAGATCCATCAGGGACCAAA 180
L N M H M N V Q N G K W D S D P S G T K
50 60
7 ACCTGCATTGATACCAAGGAAGGCATCTCGCATATTGCCAAGAAGTCTACCCCTGAAC TG 240
T C I D T K E G I L Q Y C Q E V Y P E L
70 80
8 CAGATCACCAATGTGGTAGAAGCCAACCAACCACTGACCATCCAGAACTGGTGAAGCGG 300
Q I T N V V E A N Q P V T I Q N W C K R
90 100
9 GGCCGCAAGCAGTGCAAGACCCATCCCCACTTTGTGATTCCCTACCGCTGCTTAGTGGT 360
G R K Q C K T H P H F V I P Y R C L V G
110 120
10 GAGTTTGTAAGTGATGCCCTTCTCGTTCCTGACAAGTGCAAATCTTACACCAGGAGAGG 420
E F V S D A L L V P D K C K F L H Q E R
130 140
11 ATGGATGTTTGGCAAACTCATCTTCACTGGCACACCGTCGCCAAAGAGACATGCAGTGAG 480
M D V C E T H L H W H T V A K E T C S E
150 160
12 AAGAGTACCAACTTGCATGACTACGGCATGTTGCTGCCCTGCGGAATTGACAAGTTCGGA 540
K S T N L H D Y G M L L P C G I D K F R
170 180
13 GGGGTAGAGTTTGTGTGTTGCCCACTGGCTGAAGAAAGTGACAATGTGGATTCTGCTGAT 600
G V E F V C C P L A E E S D N V D S A D
190 200
14 GCGGAGGAGGATGACTCGGATGTCTGGTGGGGCGGAGCAGACACAGACTATGCAGATGGG 660
A E E D D S D V W G G A D T D Y A D G
210 220
15 AGTGAAGACAAAGTAGTAGAAGTAGCAGAGGAGGAAGAAGTGGCTGAGGTGGAAGAAGAA 720
S E D K V V E V A E E E E V A E V E E E
230 240
16 GAAGCCGATGATGACGAGGACGATGAGGATGGTGATGAGGTAGAGGAAGAGGCTGAGGAA 780
E A D D D E D D E D G D E V E E E A E E
250 260
17 CCCTACGAAGAAGCCACAGAGAGAACCACGATTGCCACCACCACCACCACCACCACA 840
P Y E E A T E R T T S I A T T T T T T T
270 280
18 GAGTCTGTGGAAGAGGTGGTTCGAGTTCCTACAACAGCAGCCAGTACCCCTGATGCCGTT 900
E S V E E V V R V P T T A A S T P D A V
290 300
19 GACAAGTATCTCGAGACACCTGGGGATGAGAATGAACATGCCCATTTCCAGAAAGCCAAA 960
D K Y L E T P G D E N E H A H F Q K A K
310 320
20 GAGAGGCTTGAGGCCAAGCACCAGAGAGAAATGTCACGGTCATGAGAGAATGGGAAGAG 1020
E R L E A K H R E R M S Q V M R E W E E
330 340
21 GCAGAACGTCAAGCAAAGAACTTGCTAAAGCTGATAAGAAGGCAGTTATCCAGCATTTTC 1080
A E R Q A K N L P K A D K K A V I Q H F
350 360
22 CAGGAGAAAGTGAATCTTTGGAACAGGAAGCAGCCAACGAGAGACAGCAGCTGGTGGAG 1140
Q E K V E S L E Q E A A N E R Q Q L V E
370 380
23 ACACACATGGCCAGAGTGAAGCCATGCTCAATGACCGCCGCCGCTGGCCCTGGAGAAC 1200
T H M A R V E A M L N D R R R L A L E N
390 400
24 TACATCACCGCTCTGCAGGCTGTTCTCCTCGGCCTCGTCACGTGTTCAATATGCTAAAG 1260
Y I T A L Q A V P P R P R H V F N M L K
410 420
25 AAGTATGTCCGCGCAGAACAGAAGGACAGACAGCACACCCTAAAGCATTTTCGAGCATGTG 1320
C Y V R A E Q K D R Q H T L K H F E H V
430 440

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10/10
FIG. 9B

26 CGCATGGTGGATCCCAAGAAAGCCGCTCAGATCCGGTCCCAGGTTATGACACACCTCCGT 1380
 R M V D P K K A A Q I R S Q V M T M L R
 450 460

27 GTGATTTATGAGCGCATGAATCAGTCTCTCTCCCTGCTCTACAACGTGCCTGCAGTGGCC 1440
 V I Y E R M N Q S L S L L Y N V P A V A
 470 480

28 GAGGAGATTGAGGATGAAGTTGATGAGCTGCTTCAGAAAGAGCAAACTATTTCAGATGAC 1500
 E E I Q D E V D E L L Q K E Q N Y S D D
 490 500

29 GTCTTGGCCAACATGATTAGTGAACCAAGGATCAGTTACGGAACGATGCTCTCATGCCA 1560
 V L A N M I S E P R I S Y G N D A L M P
 510 520

30 TCTTTGACCGAAACGAAACACCGTGGAGCTCCTTCCCGTGAATGGAGAGTTACAGCTG 1620
 S L T E T K T T V E L L P V N G E F S L
 530 540

31 GACGATCTCCAGCCGTGGCATTCTTTTGGGGCTGACTCTGTGCCAGCCAACACAGAAAAC 1680
 D D L Q P W H S F G A D S V P A N T E N
 550 560

32 GAAGTTGAGCCTGTTGATGCCCGCCCTGCTGCCGACCGAGGACTGACCACTCGACCAGGT 1740
 E V E P V D A R P A A D R G L T T R P G
 570 580

33 TCTGGGTTGACAAATATCAAGACGGAGGAGATCTCTGAAGTGAAGATGGATGCAGAATTC 1800
 S G L T N I K T E E I S E V K M D A E F
 590 600

34 CGACATGACTCAGGATATGAAGTTCATCATCAAAAATTGGTGTCTTTGCAGAAGATGTG 1860
 R H D S G Y E V H H O K L V F F A E D V
 610 620

35 GGTTCAAACAAAGGTGCAATCATTGGACTCAGTGGTGGGCGGTGTTGTTCATAGCGACAGTG 1920
 G S N K G A I I G L M V G G V V I A I T V
 630 640

36 ATCGTCATCACCTTGCTGATGCTGAAGAAGAAACAGTACACATCCATTTCATCATGGTGTG 1980
 I V I T L V M L K K K Q Y T S I H H G V
 650 660

37 GTGGAGGTTGACGCCGTGTCACCCCAGAGGAGCGCCACCTGTCCAAGATGCAGCAGAAC 2040
 V E V D A A V T P E E R H L S K M Q Q N
 670 680

38 GGCTACGAAAATCCAACCTACAAGTTCTTTGAGCAGATGCAGAACTAGACCCCCGCCACA 2100
 G Y E N P T Y K F F E Q M Q N *
 690

39 GCAGCCTCTGAAGTTGGACAGCAAAACCATTGCTTCACTACCCATCGGTGTCCATTTTATA 2160

40 GAATAATGTGGGAAGAAACAAACCCGTTTTATGATTTACTCATTATCGCCTTTTGACAGC 2220

41 TGTGCTGTAACACAAGTAGATGCCTGAACCTGAATTAATCCACACATCAGTAATGTATTC 2280

42 TATCTCTCTTTACATTTTGGTCTCTATACTACATTATTAATGGGTTTTGTGTACTGTAAA 2340

43 GAATTTAGCTGTATCAAACCTAGTGCATGAATAGATTCTCTCCTGATTATTTATCACATAG 2400

44 CCCCTTAGCCAGTTGTATATTATCTTGTGGTTTGTGACCAATTAAGTCTACTTTTACA 2460

45 TATGCTTTAAGAATCGATGGGGGATGCTTCATGTGAACGTGGGAGTTCAGCTGCTTCTCT 2520

46 TGCCTAAGTATTCCTTTCCCTGATCACTATGCATTTTAAAGTTAAACATTTTAAAGTATTT 2580

47 CAGATGCTTTAGAGAGATTTTTCATGACTGCATTTTACTGTACAGATTGCTGCTTC 2640

48 TGCTATATTTGTGATATAGGAATTAAGAGGATACACACGTTTGTCTTTCGTGCCTGTTT 2700

49 TATGTGCACACATTAGGCATTGAGACTTCAAGCTTTTCTTTTTTTGTCCACGTATCTTTG 2760

50 GGTCTTTGATAAAGAAAAGAAATCCCTGTTCAATGTAAGCACTTTTACGGGGCGGGTGGGG 2820

51 AGGGGTGCTCTGCTGGTCTTCAATTACCAAGAATTCTCCAAAACAATTTTCTGCAGGATG 2880

52 ATTGTACAGAATCATTGCTTATGACATGATCGCTTTCTACACTGTATTACATAAATAAAT 2940

53 TAAATAAATAAACCCTGGGCAAGACTTTTCTTTGAAGGATGACTACAGACATTAAATAAT 3000

54 CGAAGTAATTTGGGTGGGGAGAAGAGGCAGATTCAATTTTCTTTAACCAGTCTGAAGTT 3060

55 TCATTTATGATACAAAAGAAGATGAAAATGGAAGTGGAATATAAGGGGATGAGGAAGGC 3120

56 ATGCCTGGACAAACCTTCTTTTAAAGATGTGTCTTCAATTTGTATAAATGGTGTTTTCA 3180

57 TGTAATAAATACATTCTTGGAGGAGC-poly (A) tail

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INTERNATIONAL SEARCH REPORT

I .national application No.
PCT/US94/05809

A. CLASSIFICATION OF SUBJECT MATTER

IPC(5) :G01N 33/367; A61K 49/00, 43/00

US CL :435/7.21; 424/1.11, 1.57, 9; 514/2

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 435/7.1, 7.2, 7.21; 424/1.11, 1.57, 1.69, 9; 514/2

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

Medline, EMBASE, BIOSIS, CA Search, WPO, APS, IntelliGenetics

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X Y	WO, A, 93/04194 (MAGGIO ET AL) 04 March 1993, see entire document, especially page 4, lines 8-15, page 8, lines 14-17, page 12, lines 15-39 and page 14, lines 26-36.	1, 2, 6, 7, 8-15 3-5
Y	Proceedings of the National Academy of Science USA, Volume 87, issued June 1992, Maggio et al, "Reversible in vitro growth of Alzheimer disease beta-amyloid plaques by deposition of labeled amyloid peptide", pages 5462-5466, especially see Abstract on page 5462.	1-15

☒ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents:	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A" document defining the general state of the art which is not considered to be of particular relevance	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
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"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"&" document member of the same patent family
"O" document referring to an oral disclosure, use, exhibition or other means	
"P" document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

24 AUGUST 1994

Date of mailing of the international search report

SEP 06 1994

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INTERNATIONAL SEARCH REPORT

International application No.
PCT/US94/05809

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X Y	Society for Neuroscience Abstracts from the 21st Annual Meeting, Volume 17, Number 1-2, issued Nov. 1991, Mantyh et al, "Distribution and Characterization of Amyloid Beta Protein Deposition in Normal Human and Alzheimer's Diseased Cerebral Cortex Using 125I-BAP as the Radioligand", page 912, Abstract 364.4, see entire abstract.	1, 2, 6, 7 3-5, 8-15
Y	Nature, Volume 325, issued 19 February 1987, Kang et al "The precursor of Alzheimer's disease amyloid A4 protein resembles a cell-surface receptor", pages 733-735, especially see Fig. 2.	1-15
A	Freeman et al, editors, Physician Desk Reference For Radiology and Nuclear Medicine, published 1977 by Medical Economics Company, Oradell, N.J., pages 39-43.	1-15
A	Proceedings of the National Academy of Science USA, Volume 88, issued August 1991, Kowall et al, "An in vivo model for the neurodegenerative effects of beta amyloid and protection by substance P", pages 7247-7251.	1-15
A	Neurobiology of Aging, Volume 13, Number 5, issued 1992, May et al, "Beta-Amyloid Peptide In Vitro Toxicity: Lot-to-Lit Variability", pages 605-607.	1-15